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## (57) Abstract

The invention disclosed in this patent document relates to transmembrane receptors, more particularly to endogenous, human orphan G protein-coupled receptors:

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## HUMAN ORPHAN G PROTEIN-COUPLED RECEPTORS

This patent document claims priority benefit of each of the following applications, all filed with the United States Patent and Trademark Office via U.S. Express Mail on the 5 indicated filing dates: U.S. Provisional Number 60/121,852, filed; February 26, 1999 claiming the benefit of U.S. Provisional Number 60/109,213, filed November 20, 1998; U.S. Provisional Number 60/120,416, filed February 16, 1999; U.S. Provisional Number 60/123,946, filed March 12, 1999; U.S. Provisional Number 60/123,949, filed March 12, 1999; U.S. Provisional Number 60/136,436, filed May 28, 1999; U.S. Provisional 10 Number 60/136,439, filed May 28, 1999; U.S. Provisional Number 60/136,567, filed May 28, 1999; U.S. Provisional Number 60/137,127, filed May 28, 1999; U.S. Provisional Number 60/137,131, filed May 28, 1999; U.S. Provisional Number 141,448, filed June 29, 1999 claiming priority from U.S. Provisional Number 60/136,437, filed May 28, 1999; U.S. Provisional Number \_\_ (Arena Pharmaceuticals, Inc. docket number 15 CHN10-1), filed September 29, 1999; U.S. Provisional Number 60/156,333, filed September 29, 1999; U.S. Provisional Number 60/156,555, filed September 29, 1999; U.S. Provisional Number 60/156,634, filed September 29, 1999; U.S. Provisional \_ (Arena Pharmaceuticals, Inc. docket number RUP6-1), filed October 1, 1999; U.S. Provisional Number \_ (Arena Pharmaceuticals, Inc. docket number 20 RUP7-1), filed October 1, 1999; U.S. Provisional Number Pharmaceuticals, Inc. docket number CHN6-1), filed October 1, 1999; U.S. Provisional

Number \_\_\_\_\_\_ (Arena Pharmaceuticals, Inc. docket number RUP5-1), filed October 1, 1999; U.S. Provisional Number \_\_\_\_\_\_ (Arena Pharmaceuticals, Inc. docket number CHN9-1), filed October 1, 1999. This patent document is related to U.S. Serial Number 09/170,496 filed October 13, 1998, and U.S. Serial Number unknown (Woodcock 5 Washburn Kurtz Mackiewicz & Norris, LLP docket number AREN-0054) filed on October 12, 1999 (via U.S. Express Mail) both being incorporated herein by reference.

This patent document also is related to U.S. Serial No. 09/364,425; filed July 30, 1999, which is incorporated by reference in its entirety. This application also claims priority to U.S. Serial Number \_\_\_\_\_ (Woodcock, Washburn, Kurtz, Makiewicz & Norris, LLP 10 docket number AREN-0050), filed on October 12, 1999 (via U.S. Express Mail), incorporated by reference herein in its entirety. Each of the foregoing applications are incorporated herein by reference in their entirety.

#### FIELD OF THE INVENTION

The invention disclosed in this patent document relates to transmembrane receptors, 15 and more particularly to endogenous, orphan, human G protein-coupled receptors ("GPCRs").

#### BACKGROUND OF THE INVENTION

Although a number of receptor classes exist in humans, by far the most abundant and therapeutically relevant is represented by the G protein-coupled receptor (GPCR or GPCRs) 20 class. It is estimated that there are some 100,000 genes within the human genome, and of these, approximately 2% or 2,000 genes, are estimated to code for GPCRs. Receptors, including GPCRs, for which the endogenous ligand has been identified are referred to as "known" receptors, while receptors for which the endogenous ligand has not been identified

are referred to as "orphan" receptors. GPCRs represent an important area for the development of pharmaceutical products: from approximately 20 of the 100 known GPCRs, 60% of all prescription pharmaceuticals have been developed. This distinction is not merely semantic, particularly in the case of GPCRs. Thus, the orphan GPCRs are to the 5 pharmaceutical industry what gold was to California in the late 19th century – an opportunity to drive growth, expansion, enhancement and development.

GPCRs share a common structural motif. All these receptors have seven sequences of between 22 to 24 hydrophobic amino acids that form seven alpha helices, each of which spans the membrane (each span is identified by number, *i.e.*, transmembrane-1 (TM-1), 10 transmebrane-2 (TM-2), etc.). The transmembrane helices are joined by strands of amino acids between transmembrane-2 and transmembrane-3, transmembrane-4 and transmembrane-5, and transmembrane-6 and transmembrane-7 on the exterior, or "extracellular" side, of the cell membrane (these are referred to as "extracellular" regions 1, 2 and 3 (EC-1, EC-2 and EC-3), respectively). The transmembrane helices are also joined 15 by strands of amino acids between transmembrane-1 and transmembrane-2, transmembrane-3 and transmembrane-4, and transmembrane-5 and transmembrane-6 on the interior, or "intracellular" side, of the cell membrane (these are referred to as "intracellular" regions 1, 2 and 3 (IC-1, IC-2 and IC-3), respectively). The "carboxy" ("C") terminus of the receptor lies in the intracellular space within the cell, and the "amino" ("N") terminus of the receptor 20 lies in the extracellular space outside of the cell.

Generally, when an endogenous ligand binds with the receptor (often referred to as "activation" of the receptor), there is a change in the conformation of the intracellular region that allows for coupling between the intracellular region and an intracellular "G-protein." It

has been reported that GPCRs are "promiscuous" with respect to G proteins, i.e., that a GPCR can interact with more than one G protein. See, Kenakin, T., 43 Life Sciences 1095 (1988). Although other G proteins exist, currently, Gq, Gs, Gi, and Go are G proteins that have been identified. Endogenous ligand-activated GPCR coupling with the G-protein begins a signaling cascade process (referred to as "signal transduction"). Under normal conditions, signal transduction ultimately results in cellular activation or cellular inhibition. It is thought that the IC-3 loop as well as the carboxy terminus of the receptor interact with the G protein.

Under physiological conditions, GPCRs exist in the cell membrane in equilibrium 10 between two different conformations: an "inactive" state and an "active" state. A receptor in an inactive state is unable to link to the intracellular signaling transduction pathway to produce a biological response. Changing the receptor conformation to the active state allows linkage to the transduction pathway (via the G-protein) and produces a biological response. A receptor may be stabilized in an active state by an endogenous ligand or a compound such 15 as a drug.

#### SUMMARY OF THE INVENTION

Disclosed herein are human endogenous orphan G protein-coupled receptors.

#### BRIEF DESCRIPTION OF THE DRAWINGS

Figures 1A and 1B provide reference "grids" for certain dot-blots provided herein 20 (see also, Figure 2A and 2B, respectively).

Figures 2A and 2B provide reproductions of the results of certain dot-blot analyses resulting from hCHN3 and hCHN8, respectively (see also, Figures 1A and 1B, respectively).

Figure 3 provides a reproduction of the results of RT-PCR analysis of hRUP3.

Figure 4 provides a reproduction of the results of RT-PCR analysis of hRUP4.

Figure 5 provides a reproduction of the results of RT-PCR analysis of hRUP6.

# DETAILED DESCRIPTION

The scientific literature that has evolved around receptors has adopted a number of 5 terms to refer to ligands having various effects on receptors. For clarity and consistency, the following definitions will be used throughout this patent document. To the extent that these definitions conflict with other definitions for these terms, the following definitions shall control:

AMINO ACID ABBREVIATIONS used herein are set out in Table 1:

10		TABLE 1		
	ALANINE	ALA	A	
	ARGININE	ARG	R	
	ASPARAGINE	ASN	N	
•	ASPARTIC ACID	ASP	D.	
15	CYSTEINE	CYS	ć	
	GLUTAMIC ACID	GLU	E	
	GLUTAMINE	GLN	ō	
	GLYCINE	GLY	G ·	•
	HISTIDINE	HIS	H	
20	ISOLEUCINE	ILE	ī	
	LEUCINE	LEU		
	LYSINE	LYS	$\tilde{\mathbf{K}}$	
	METHIONINE	MET	M	•
	PHENYLALANINE	PHE	· · · · · · · · · · · · · · · · · · ·	٠.
25	PROLINE	PRO	P	,
	SERINE	SER	Š	7
	THREONINE	THR	Ť	* .
	TRYPTOPHAN	TRP	w	
	TYROSINE	TYR	Ÿ	
30	VALINE	VAL	v	

COMPOSITION means a material comprising at least one component.

ENDOGENOUS shall mean a material that a mammal naturally produces. ENDOGENOUS in reference to, for example and not limitation, the term "receptor," shall mean that which is naturally produced by a mammal (for example, and not limitation, a

human) or a virus. By contrast, the term NON-ENDOGENOUS in this context shall mean that which is not naturally produced by a mammal (for example, and not limitation, a human) or a virus.

HOST CELL shall mean a cell capable of having a Plasmid and/or Vector 5 incorporated therein. In the case of a prokaryotic Host Cell, a Plasmid is typically replicated as a autonomous molecule as the Host Cell replicates (generally, the Plasmid is thereafter isolated for introduction into a eukaryotic Host Cell); in the case of a eukaryotic Host Cell, a Plasmid is integrated into the cellular DNA of the Host Cell such that when the eukaryotic Host Cell replicates, the Plasmid replicates. Preferably, for the purposes of the invention 10 disclosed herein, the Host Cell is eukaryotic, more preferably, mammalian, and most preferably selected from the group consisting of 293, 293T and COS-7 cells.

LIGAND shall mean an endogenous, naturally occurring molecule specific for an endogenous, naturally occurring receptor.

NON-ORPHAN RECEPTOR shall mean an endogenous naturally occurring 15 molecule specific for an endogenous naturally occurring ligand wherein the binding of a ligand to a receptor activates an intracellular signaling pathway.

ORPHAN RECEPTOR shall mean an endogenous receptor for which the endogenous ligand specific for that receptor has not been identified or is not known.

PLASMID shall mean the combination of a Vector and cDNA. Generally, a Plasmid 20 is introduced into a Host Cell for the purposes of replication and/or expression of the cDNA as a protein.

VECTOR sin reference to cDNA shall mean a circular DNA capable of incorporating at least one cDNA and capable of incorporation into a Host Cell.

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The order of the following sections is set forth for presentational efficiency and is not intended, nor should be construed, as a limitation on the disclosure or the claims to follow.

## Identification of Human GPCRs

of information regarding nucleic acid sequences located within the human genome; it has been the case in this endeavor that genetic sequence information has been made available without an understanding or recognition as to whether or not any particular genomic sequence does or may contain open-reading frame information that translate human proteins.

10 Several methods of identifying nucleic acid sequences within the human genome are within the purview of those having ordinary skill in the art. For example, and not limitation, a variety of GPCRs, disclosed herein, were discovered by reviewing the GenBank<sup>TM</sup> database, while other GPCRs were discovered by utilizing a nucleic acid sequence of a GPCR, previously sequenced, to conduct a BLAST<sup>TM</sup> search of the EST database. Table A, below, 15 lists the disclosed endogenous orphan GPCRs along with a GPCR's respective homologous GPCR:

I ADLE A

	Disclosed	Accession	Open Reading	Per Cent	Reference To
	Human	Number	Frame	Homology	Homologous
20	Orphan	Identified	(Base Pairs)	To Designated	GPCR
•	<b>GPCRs</b>			GPCR	(Accession No.)
,	·•	. **			
	hARE-3 hARE-4	AL033379 AC006087	1,260 bp 1,119 bp	52.3% LPA-R 36% P2Y5	U92642 AF000546

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	hARE-5	AC006255	1,104 bp	32% Oryzias	D43633
	CDDCC			latipes	
	hGPR27	AA775870	1,128 bp		
	hARE-1	A1090920	999 bp	43%	D13626
				VIA 40001	
	hARE-2	AA359504	1,122 bp	KIAA0001	
5	hPPR1	H67224	1,053 bp	53% GPR27	7.01.504
	hG2A	AA754702	1,033 bp	39% EBI1	L31581
	hRUP3	AL035423		31% GPR4	L36148
		712033423	1,005 bp	30%	2133653
				Drosophila	
	hRUP4	41202660		melanogaster	
	nRUP4	AI307658	1,296 bp	32% pNPGPR	NP_004876
				28% and 29 %	AAC41276
					7111041270
				Zebra fish Ya	and
				and Yb,	
					AAB94616
				respectively	
	hRUP5	AC005849	1,413 bp	25% DEZ	Q99788
10	hRUP6	4.0005024		23% FMLPR	P21462
10	hRUP7	AC005871	1,245 bp	48% GPR66	NP_006047
	hCHN3	AC007922	1,173 bp	43% H3R	AF140538
	hCHN4	EST 36581	1,113 bp	53% GPR27	
	hCHN6	AA804531	1,077 bp	32% thrombin	4503637
15		EST 2134670	1,503 bp	36% edg-1	NP_001391
13	hCHN8	EST 764455	1,029 bp	47%	D13626
101	hCHN9	EST 1541536	1.075	KIAA0001	<b>₩</b>
	hCHN10	EST 1365839	1,077 bp	41% LTB4R	NM_000752
	IICIII(10	E91 1303839	1,055 bp	35% P2Y	NM_002563

Receptor homology is useful in terms of gaining an appreciation of a role of the disclosed receptors within the human body. Additionally, such homology can provide insight 20 as to possible endogenous ligand(s) that may be natural activators for the disclosed orphan GPCRs.

## B. Receptor Screening

Techniques have become more readily available over the past few years for

endogenous-ligand identification (this, primarily, for the purpose of providing a means of conducting receptor-binding assays that require a receptor's endogenous ligand) because the traditional study of receptors has always proceeded from the a priori assumption (historically based) that the endogenous ligand must first be identified before discovery could proceed to 5 find antagonists and other molecules that could affect the receptor. Even in cases where an antagonist might have been known first, the search immediately extended to looking for the endogenous ligand. This mode of thinking has persisted in receptor research even after the discovery of constitutively activated receptors. What has not been heretofore recognized is that it is the active state of the receptor that is most useful for discovering agonists, partial 10 agonists, and inverse agonists of the receptor. For those diseases which result from an overly active receptor or an under-active receptor, what is desired in a therapeutic drug is a compound which acts to diminish the active state of a receptor or enhance the activity of the receptor, respectively, not necessarily a drug which is an antagonist to the endogenous ligand. This is because a compound that reduces or enhances the activity of the active receptor state 15 need not bind at the same site as the endogenous ligand. Thus, as taught by a method of this invention, any search for therapeutic compounds should start by screening compounds against the ligand-independent active state.

As is known in the art, GPCRs can be "active" in their endogenous state even without the binding of the receptor's endogenous ligand thereto. Such naturally-active receptors can 20 be screened for the direct identification (i.e., without the need for the receptor's endogenous ligand) of, in particular, inverse agonists. Alternatively, the receptor can be "activated" via, e.g., mutation of the receptor to establish a non-endogenous version of the receptor that is active in the absence of the receptor's endogenous ligand.

Screening candidate compounds against an endogenous or non-endogenous, constitutively activated version of the human orphan GPCRs disclosed herein can provide for the direct identification of candidate compounds which act at this cell surface receptor, without requiring use of the receptor's endogenous ligand. By determining areas within 5 the body where the endogenous version of human GPCRs disclosed herein is expressed and/or over-expressed, it is possible to determine related disease/disorder states which are associated with the expression and/or over-expression of the receptor; such an approach is disclosed in this patent document.

With respect to creation of a mutation that may evidence constitutive activation of 10 human orphan GPCRs disclosed herein is based upon the distance from the proline residue at which is presumed to be located within TM6 of the GPCR typically nears the TM6/IC3 interface (such proline residue appears to be quite conserved). By mutating the amino acid residue located 16 amino acid residues from this residue (presumably located in the IC3 region of the receptor) to, most preferably, a lysine residue, such activation may be obtained. 15 Other amino acid residues may be useful in the mutation at this position to achieve this objective.

#### C. Disease/Disorder Identification and/or Selection

Preferably, the DNA sequence of the human orphan GPCR can be used to make a probe for (a) dot-blot analysis against tissue-mRNA, and/or (b) RT-PCR identification of 20 the expression of the receptor in tissue samples. The presence of a receptor in a tissue source, or a diseased tissue, or the presence of the receptor at elevated concentrations in diseased tissue compared to a normal tissue, can be preferably utilized to identify a correlation with a treatment regimen, including but not limited to, a disease associated

with that disease. Receptors can equally well be localized to regions of organs by this technique. Based on the known functions of the specific tissues to which the receptor is localized, the putative functional role of the receptor can be deduced.

# D. Screening of Candidate Compounds

## 1. Generic GPCR screening assay techniques

When a G protein receptor becomes constitutively active (i.e., active in the absence of endogenous ligand binding thereto), it binds to a G protein (e.g., Gq, Gs, Gi, Go) and stimulates the binding of GTP to the G protein. The G protein then acts as a GTPase and slowly hydrolyzes the GTP to GDP, whereby the receptor, under normal conditions, becomes 10 deactivated. However, constitutively activated receptors continue to exchange GDP to GTP. A non-hydrolyzable analog of GTP, [35S]GTPγS, can be used to monitor enhanced binding to membranes which express constitutively activated receptors. It is reported that [35S]GTPγS can be used to monitor G protein coupling to membranes in the absence and presence of ligand. An example of this monitoring, among other examples well-known and 15 available to those in the art, was reported by Traynor and Nahorski in 1995. The preferred use of this assay system is for initial screening of candidate compounds because the system is generically applicable to all G protein-coupled receptors regardless of the particular G protein that interacts with the intracellular domain of the receptor.

# 2. Specific GPCR screening assay techniques

Once candidate compounds are identified using the "generic" G protein-coupled receptor assay (i.e., an assay to select compounds that are agonists, partial agonists, or inverse agonists), further screening to confirm that the compounds have interacted at the receptor site is preferred. For example, a compound identified by the "generic" assay may not bind to the

receptor, but may instead merely "uncouple" the G protein from the intracellular domain.

#### a. Gs and Gi.

Gs stimulates the enzyme adenylyl cyclase. Gi (and Go), on the other hand, inhibit Adenylyl cyclase catalyzes the conversion of ATP to cAMP; thus, 5 constitutively activated GPCRs that couple the Gs protein are associated with increased cellular levels of cAMP. On the other hand, constitutively activated GPCRs that couple the Gi (or Go) protein are associated with decreased cellular levels of cAMP. See, generally, "Indirect Mechanisms of Synaptic Transmission," Chpt. 8, From Neuron To Brain (3rd Ed.) Nichols, J.G. et al eds. Sinauer Associates, Inc. (1992). Thus, assays that detect cAMP can 10 be utilized to determine if a candidate compound is, e.g., an inverse agonist to the receptor (i.e., such a compound would decrease the levels of cAMP). A variety of approaches known in the art for measuring cAMP can be utilized; a most preferred approach relies upon the use of anti-cAMP antibodies in an ELISA-based format. Another type of assay that can be utilized is a whole cell second messenger reporter system assay. Promoters on genes drive 15 the expression of the proteins that a particular gene encodes. Cyclic AMP drives gene expression by promoting the binding of a cAMP-responsive DNA binding protein or transcription factor (CREB) which then binds to the promoter at specific sites called cAMP response elements and drives the expression of the gene. Reporter systems can be constructed which have a promoter containing multiple cAMP response elements before the reporter 20 gene, e.g., B-galactosidase or luciferase. Thus, a constitutively activated Gs-linked receptor causes the accumulation of cAMP that then activates the gene and expression of the reporter protein. The reporter protein such as β-galactosidase or luciferase can then be detected using standard biochemical assays (Chen et al. 1995).

## Go and Gq.

Gq and Go are associated with activation of the enzyme phospholipase C, which in turn hydrolyzes the phospholipid PIP2, releasing two intracellular messengers: 5 diacycloglycerol (DAG) and inistol 1,4,5-triphoisphate (IP3). Increased accumulation of IP3 is associated with activation of Gq- and Go-associated receptors. See, generally, "Indirect Mechanisms of Synaptic Transmission," Chpt. 8, From Neuron To Brain (3rd Ed.) Nichols, J.G. et al eds. Sinauer Associates, Inc. (1992). Assays that detect IP3 accumulation can be utilized to determine if a candidate compound is, e.g., an inverse agonist to a Gq- or Go-10 associated receptor (i.e., such a compound would decrease the levels of IP3). Gq-associated receptors can also been examined using an AP1 reporter assay in that Gq-dependent phospholipase C causes activation of genes containing AP1 elements; thus, activated Gq-associated receptors will evidence an increase in the expression of such genes, whereby inverse agonists thereto will evidence a decrease in such expression, and agonists will 15 evidence an increase in such expression. Commercially available assays for such detection are available.

## 3. GPCR Fusion Protein

The use of an endogenous, constitutively activated orphan GPCR, or a non-endogenous, constitutively activated orphan GPCR, for screening of candidate compounds 20 for the direct identification of inverse agonists, agonists and partial agonists provides a unique challenge in that, by definition, the receptor is active even in the absence of an endogenous ligand bound thereto. Thus, it is often useful that an approach be utilized that can enhance the signal obtained by the activated receptor. A preferred approach is the use of a GPCR Fusion Protein.

Generally, once it is determined that a GPCR is or has been constitutively activated, using the assay techniques set forth above (as well as others), it is possible to determine the predominant G protein that couples with the endogenous GPCR. Coupling of the G protein to the GPCR provides a signaling pathway that can be assessed. Because it is most preferred 5 that screening take place by use of a mammalian expression system, such a system will be expected to have endogenous G protein therein. Thus, by definition, in such a system, the constitutively activated orphan GPCR will continuously signal. In this regard, it is preferred that this signal be enhanced such that in the presence of, e.g., an inverse agonist to the receptor, it is more likely that it will be able to more readily differentiate, particularly in the 10 context of screening, between the receptor when it is contacted with the inverse agonist.

The GPCR Fusion Protein is intended to enhance the efficacy of G protein coupling with the GPCR. The GPCR Fusion Protein is preferred for screening with a non-endogenous, constitutively activated GPCR because such an approach increases the signal that is most preferably utilized in such screening techniques, although the GPCR Fusion 15 Protein can also be (and preferably is) used with an endogenous, constitutively activated GPCR. This is important in facilitating a significant "signal to noise" ratio; such a significant ratio is import preferred for the screening of candidate compounds as disclosed herein.

The construction of a construct useful for expression of a GPCR Fusion Protein is within the purview of those having ordinary skill in the art. Commercially available 20 expression vectors and systems offer a variety of approaches that can fit the particular needs of an investigator. The criteria of importance for such a GPCR Fusion Protein construct is that the GPCR sequence and the G protein sequence both be in-frame (preferably, the sequence for the GPCR is upstream of the G protein sequence) and that the "stop" codon of

the GPCR must be deleted or replaced such that upon expression of the GPCR, the G protein can also be expressed. The GPCR can be linked directly to the G protein, or there can be spacer residues between the two (preferably, no more than about 12, although this number can be readily ascertained by one of ordinary skill in the art). We have a preference (based 5 upon convenience) of use of a spacer in that some restriction sites that are not used will, effectively, upon expression, become a spacer. Most preferably, the G protein that couples to the GPCR will have been identified prior to the creation of the GPCR Fusion Protein construct. Because there are only a few G proteins that have been identified, it is preferred that a construct comprising the sequence of the G protein (i.e., a universal G protein 10 construct) be available for insertion of an endogenous GPCR sequence therein; this provides for efficiency in the context of large-scale screening of a variety of different endogenous GPCRs having different sequences.

## E. Other Utility

Although a preferred use of the human orphan GPCRs disclosed herein may be for 15 the direct identification of candidate compounds as inverse agonists, agonists or partial agonists (preferably for use as pharmaceutical agents), these versions of human GPCRs can also be utilized in research settings. For example, in vitro and in vivo systems incorporating GPCRs can be utilized to further elucidate and understand the roles these receptors play in the human condition, both normal and diseased, as well as understanding the role of 20 constitutive activation as it applies to understanding the signaling cascade. The value in human orphan GPCRs is that its utility as a research tool is enhanced in that by determining the location(s) of such receptors within the body, the GPCRs can be used to understand the role of these receptors in the human body before the endogenous ligand therefor is identified.

Other uses of the disclosed receptors will become apparent to those in the art based upon, inter alia, a review of this patent document.

#### **EXAMPLES**

The following examples are presented for purposes of elucidation, and not limitation, 5 of the present invention. While specific nucleic acid and amino acid sequences are disclosed herein, those of ordinary skill in the art are credited with the ability to make minor modifications to these sequences while achieving the same or substantially similar results reported below. Unless otherwise indicated below, all nucleic acid sequences for the disclosed endogenous orphan human GPCRs have been sequenced and verified. For 10 purposes of equivalent receptors, those of ordinary skill in the art will readily appreciate that conservative substitutions can be made to the disclosed sequences to obtain a functionally equivalent receptor.

# Example 1 ENDOGENOUS HUMAN GPCRS

## 1. Identification of Human GPCRs

Several of the disclosed endogenous human GPCRs were identified based upon a review of the GenBank database information. While searching the database, the following cDNA clones were identified as evidenced below.

	Disclosed	Accession	Complete DNA	Open Reading	Nucleic Acid	Amino
20	Human	Number	Sequence	Frame	SEQ.ID.	Acid
,	Orphan		(Base Pairs)	(Base Pairs)	NO.	SEQ.ID.
	GPCRs					NO.

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-	_		
1	7		
	•	•	

.•	hARE-3	AL033379	111,389 bp 1,260 bp 1 2
	hARE-4	AC006087	226,925 bp 1,119 bp 3 4
	hARE-5	AC006255	127,605 bp 1,104 bp 5 6
	hRUP3	AL035423	140,094 bp 1,005 bp 7 8
5	hRUP5	AC005849	169,144 bp 1,413 bp 9 10
	hRUP6	AC005871	218,807 bp 1,245 bp 11 12
,	hRUP7	AC007922	158,858 bp 1,173 bp 13 14

Other disclosed endogenous human GPCRs were identified by conducting a BLAST search of EST database (dbest) using the following EST clones as query sequences. The 10 following EST clones identified were then used as a probe to screen a human genomic library.

() •	Disclosed	Query	EST Clone/ Open		Nucleic Acid	Amino Acid	
	Human	(Sequence)	Accession No.	Reading	SEQ.ID.NO.	SEQ.ID.NO.	
	Orphan		Identified	Frame	,	*	
15	GPCRs hGPCR27	Mouse	AA775870	(Base Pairs) 1,125 bp	15	16	
	hARE-1	GPCR27 TDAG	1689643	999 bp	17	18	
	hARE-2	GPCR27	A1090920 68530	1,122 bp	19	20	
•	hPPR1	Bovine	AA359504 238667	1,053 bp	21	22	
20	hG2A	PPR1 Mouse	H67224 See Example 2(a),	1,113 bp	23	24	
· ·		1179426	below			* 4	

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		- 18 -			
hCHN3	N.A.	EST 36581	1,113 bp	25	26
		(full length)			
hCHN4	TDAG	1184934	1,077 bp	27	28
		AA804531			
hCHN6	N.A.	EST 2134670	1,503 bp	29	30
		(full length)			
hCHN8	KIAA0001	EST 764455	1,029 bp	31	32
hCHN 9	1365839	EST 1541536	1,077 bp	33 -	34
hCHN10	Mouse EST	Human 1365839	1,005 bp	35	36
	1365839				
hRUP4	N.A.	AI307658	1,296 bp	37	38
	N.A. = "not ap	oplicable".			

## 2. Full Length Cloning

## a. hG2A (Seq. Id. Nos. 23 & 24)

Mouse EST clone 1179426 was used to obtain a human genomic clone containing all but three amino acid hG2A coding sequences. The 5'end of this coding sequence was obtained by using 5'RACE™, and the template for PCR was Clontech's Human Spleen Marathon-ready™ cDNA. The disclosed human G2A was amplified by PCR using the G2A 15 cDNA specific primers for the first and second round PCR as shown in SEQ.ID.NO.: 39 and SEQ.ID.NO.:40 as follows:

5'-CTGTGTACAGCAGTTCGCAGAGTG-3' (SEQ.ID.NO.: 39; 1" round PCR)

5'-GAGTGCCAGGCAGAGCAGGTAGAC-3' (SEQ.ID.NO.: 40; second round PCR).

PCR was performed using Advantage™ GC Polymerase Kit (Clontech; manufacturing 20 instructions will be followed), at 94°C for 30 sec followed by 5 cycles of 94°C for 5 sec and 72°C for 4 min; and 30 cycles of 94° for 5 sec and 70° for 4 min. An approximate 1.3 Kb PCR fragment was purified from agarose gel, digested with Hind III and Xba I and cloned into the expression vector pRC/CMV2 (Invitrogen). The cloned-insert was sequenced using the T7 Sequenase™ kit (USB Amersham; manufacturer instructions will be followed) and

the sequence was compared with the presented sequence. Expression of the human G2A will be detected by probing an RNA dot blot (Clontech; manufacturer instructions will be followed) with the P<sup>32</sup>-labeled fragment.

## b. hCHN9 (Seq. Id. Nos. 33 & 34)

- Sequencing of the EST clone 1541536 indicated that hCHN9 is a partial cDNA clone having only an initiation codon; *i.e.*, the termination codon was missing. When hCHN9 was used to "blast" against the data base (nr), the 3' sequence of hCHN9 was 100% homologous to the 5' untranslated region of the leukotriene B4 receptor cDNA, which contained a termination codon in the frame with hCHN9 coding sequence. To 10 determine whether the 5' untranslated region of LTB4R cDNA was the 3' sequence of hCHN9, PCR was performed using primers based upon the 5' sequence flanking the initiation codon found in hCHN9 and the 3' sequence around the termination codon found in the LTB4R 5' untranslated region. The 5' primer sequence utilized was as follows: 5'-CCCGAATTCCTGCTTCCCCAGCTTGGCCC-3' (SEQ.ID.NO.: 41; sense) and
- PCR was performed using thymus cDNA as a template and rTth polymerase (Perkin Elmer) with the buffer system provided by the manufacturer, 0.25 uM of each primer, and 0.2 mM of each 4 nucleotides. The cycle condition was 30 cycles of 94°C for 1 min, 65°C for 1 min and 72 °C for 1 min and 10 sec. A 1.1kb fragment consistent with the predicted size was 20 obtained from PCR. This PCR fragment was subcloned into pCMV (see below) and sequenced (see, SEQ.ID.NO.: 33).

15 5'-TGTGGATCCTGTCAAAGGTCCCATTCCGG-3' (SEQ.ID.NO.: 42; antisense).

## c. hRUP 4 (Seq. Id. Nos. 37 & 38)

The full length hRUP4 was cloned by RT-PCR with human brain cDNA (Clontech)

as templates:

5'-TCACAATGCTAGGTGTGGTC-3' (SEQ.ID.NO.: 43; sense) and

5'-TGCATAGACAATGGGATTACAG-3' (SEQ.ID.NO.: 44; antisense).

PCR was performed using TaqPlus™ Precision™ polymerase (Stratagene; manufacturing 5 instructions will be followed) by the following cycles: 94°C for 2 min; 94°C 30 sec; 55°C for 30 sec, 72°C for 45 sec, and 72°C for 10 min. Cycles 2 through 4 were repeated 30 times.

The PCR products were separated on a 1% agarose gel and a 500 bp PCR fragment was isolated and cloned into the pCRII-TOPO vector (Invitrogen) and sequenced using the 10 T7 DNA Sequenase™ kit (Amsham) and the SP6/T7 primers (Stratagene). Sequence analysis revealed that the PCR fragment was indeed an alternatively spliced form of AI307658 having a continuous open reading frame with similarity to other GPCRs. The completed sequence of this PCR fragment was as follows:

5'-TCACAATGCTAGGTGTGGTCTGGCTGGTGGCAGTCATCGTAGGATCACCCATGTGGCAC
15 GTGCAACAACTTGAGATCAAATATGACTTCCTATATGAAAAGGAACACATCTGCTGCTTAGAA
GAGTGGACCAGCCCTGTGCACCAGAAGATCTACACCACCTTCATCCTTGTCATCCTCTCC
TGCCTCTTATGGTGATGCTTATTCTGTACGTAAAATTGGTTATGAACTTTGGATAAAGAAAAGA
GTTGGGGATGGTTCAGTGCTTCGAACTATTCATGGAAAAGAAATGTCCAAAATAGCCAGGAAG
AAGAAACGAGCTGTCATTATGATGGTGACAGTGGTGGCTCTCTTTGCTGTGCTGGGCACCA
20 TTCCATGTTGTCCATATGATGATTGAATACAGTAATTTTGAAAAAGGAATATGATGATGTCACA
ATCAAGATGATTTTTGCTATCGTGCAAATTATTGGATTTTCCAACTCCATCTGTAATCCCATTG
TCTATGCA-3' (SEQ.ID.NO.: 45)

Based on the above sequence, two sense oligonucleotide primer sets:

5'-CTGCTTAGAAGAGTGGACCAG-3' (SEQ.ID.NO.: 46; oligo 1),

25 5'-CTGTGCACCAGAAGATCTACAC-3' (SEQ.IDNO.: 47; oligo 2)

and two antisense oligonucleotide primer sets:

5'-CAAGGATGAAGGTGGTGTAGA-3' (SEQ.ID.NO.: 48; oligo 3)

5'-GTGTAGATCTTCTGGTGCACAGG-3' (SEQ.ID.NO.: 49; oligo 4)

were used for 3'- and 5'-race PCR with a human brain Marathon-Ready™ cDNA (Clontech,

Cat# 7400-1) as template, according to manufacture's instructions. DNA fragments generated by the RACE PCR were cloned into the pCRII-TOPO™ vector (Invitrogen) and sequenced using the SP6/T7 primers (Stratagene) and some internal primers. The 3' RACE product contained a poly(A) tail and a completed open reading frame ending at a TAA stop 5 codon. The 5' RACE product contained an incomplete 5' end; i.e., the ATG initiation codon was not present.

Based on the new 5' sequence, oligo 3 and the following primer:

5'-GCAATGCAGGTCATAGTGAGC -3' (SEQ.ID.NO.: 50; oligo 5)

were used for the second round of 5' RACE PCR and the PCR products were analyzed as 10 above. A third round of 5' RACE PCR was carried out utilizing antisense primers:

5'-TGGAGCATGGTGACGGGAATGCAGAAG-3' (SEQ.ID.NO.: 51; oligo 6) and

5'-GTGATGAGCAGGTCACTGAGCGCCAAG-3' (SEQ.ID.NO.: 52; oligo7).

The sequence of the 5' RACE PCR products revealed the presence of the initiation codon ATG, and further round of 5' RACE PCR did not generate any more 5' sequence. The 15 completed 5' sequence was confirmed by RT-PCR using sense primer

5'-GCAATGCAGGCGCTTAACATTAC-3' (SEQ.ID.NO.: 53; oligo 8)

and oligo 4 as primers and sequence analysis of the 650 bp PCR product generated from human brain and heart cDNA templates (Clontech, Cat# 7404-1). The completed 3' sequence was confirmed by RT-PCR using oligo 2 and the following antisense primer:

20 5'-TTGGGTTACAATCTGAAGGGCA-3' (SEQ.ID.NO.: 54; oligo 9)

and sequence analysis of the 670 bp PCR product generated from human brain and heart cDNA templates. (Clontech, Cat# 7404-1).

d. hRUP5 (Seq. Id. Nos. 9 & 10)

The full length hRUP5 was cloned by RT-PCR using a sense primer upstream from

ATG, the initiation codon (SEQ.ID.NO.: 55), and an antisense primer containing TCA as the stop codon (SEQ.ID.NO.: 56), which had the following sequences:

- 5'-ACTCCGTGTCCAGCAGGACTCTG-3' (SEQ.ID.NO.:55)
- 5'-TGCGTGTTCCTGGACCCTCACGTG-3' (SEQ.ID.NO.: 56)
- 5 and human peripheral leukocyte cDNA (Clontech) as a template. Advantage cDNA polymerase (Clontech) was used for the amplification in a 50ul reaction by the following cycle with step 2 through step 4 repeated 30 times: 94°C for 30 sec; 94° for 15 sec; 69° for 40 sec; 72°C for 3 min; and 72°C fro 6 min. A 1.4kb PCR fragment was isolated and cloned with the pCRII-TOPO™ vector (Invitrogen) and completely sequenced using the T7 DNA 10 Sequenase™ kit (Amsham). See, SEQ.ID.NO.: 9.

## e. hRUP6 (Seq. Id. Nos. 11 & 12)

The full length hRUP6 was cloned by RT-PCR using primers:

- 5'-CAGGCCTTGGATTTTAATGTCAGGGATGG-3' (SEQ.ID.NO.: 57) and
- 5'-GGAGAGTCAGCTCTGAAAGAATTCAGG-3' (SEQ.ID.NO.: 58);
- 15 and human thymus Marathon-Ready™ cDNA (Clontech) as a template. Advantage cDNA polymerase (Clontech, according to manufacturer's instructions) was used for the amplification in a 50ul reaction by the following cycle: 94°C for 30sec; 94°C for 5 sec; 66°C for 40sec; 72°C for 2.5 sec and 72°C for 7 min. Cycles 2 through 4 were repeated 30 times. A 1.3 Kb PCR fragment was isolated and cloned into the pCRII-TOPO™ vector (Invitrogen) 20 and completely sequenced (see, SEQ.ID.NO.: 11) using the ABI Big Dye Terminator™ kit (P.E. Biosystem).

## f. hRUP7 (Seq. Id. Nos. 13 & 14)

The full length RUP7 was cloned by RT-PCR using primers:

5'-TGATGTGATGCCAGATACTAATAGCAC-3' (SEQ.ID.NO.: 59; sense) and

5'-CCTGATTCATTTAGGTGAGATTGAGAC-3' (SEQ.ID.NO.: 60; antisense)

and human peripheral leukocyte cDNA (Clontech) as a template. Advantage™ cDNA polymerase (Clontech) was used for the amplification in a 50 ul reaction by the following cycle with step 2 to step 4 repeated 30 times: 94°C for 2 minutes; 94°C for 15 seconds; 60°C for 20 seconds; 72°C for 2 minutes; 72°C for 10 minutes. A 1.25 Kb PCR fragment was isolated and cloned into the pCRII-TOPO™ vector (Invitrogen) and completely sequenced using the ABI Big Dye Terminator™ kit (P.E. Biosystem). See, SEQ.ID.NO.: 13.

## g. hARE-5 (Seq. Id Nos. 5 & 6)

The full length hARE-5 was cloned by PCR using the hARE5 specific primers 10 5'-CAGCGCAGGGTGAAGCCTGAGAGC-3' SEQ.ID.NO.: 69 (sense, 5' of initiation codon ATG) and 5'-GGCACCTGCTGTGACCTGTGCAGG-3' SEQ.ID.NO.:70 (antisense, 3' of stop codon TGA) and human genomic DNA as template. TaqPlus Precision™ DNA polymerase (Stratagene) was used for the amplification by the following cycle with step 2 to step 4 repeated 35 times: 96°C, 2 minutes; 96°C, 20 seconds; 58°C, 30 seconds; 72°C, 2 minutes; and 72°C, 10 minutes

A 1.1 Kb PCR fragment of predicated size was isolated and cloned into the pCRII-TOPO™ vector (Invitrogen) and completely sequenced (SEQ.ID.NO.:5) using the T7 DNA Sequenase™ kit (Amsham).

## h. hARE-4 (Seq. Id. Nos.: 3 & 4)

The full length hARE-4 was cloned by PCR using the hARE-4 specific primers 5'20 CTGGTGTGCTCCATGGCATCCC-3' SEQ.ID.NO.:67 (sense, 5' of initiation codon ATG) and 5'GTAAGCCTCCCAGAACGAGAGG-3' SEQ.ID.NO.: 68 (antisense, 3' of stop codon TGA) and
human genomic DNA as template. Taq DNA polymerase (Stratagene) and 5% DMSO was
used for the amplification by the following cycle with step 2 to step 3 repeated 35 times:

94°C, 3 minutes; 94°C, 30 seconds; 59°C, 2 minutes; 72°C, 10 minutes

A 1.12 Kb PCR fragment of predicated size was isolated and cloned into the pCRII-TOPOTM-vector (Invitrogen) and completely sequenced (SEQ.ID.NO.:3) using the T7 DNA Sequenase<sup>TM</sup> kit (Amsham).

## i. hARE-3 (Seq.Id.Nos.: 1 & 2)

The full length hARE-3 was cloned by PCR using the hARE-3 specific primers 5'gatcaagcttCCATCCTACTGAAACCATGGTC-3' SEQ.ID.NO.:65 (sense, lower case nucleotides
represent Hind III overhang, ATG as initiation codon) and 5'gatcagatctCAGTTCCAATATTCACACCACCGTC-3' SEQ.ID.NO.:66 (antisense, lower case
10 nucleotides represent Xba I overhang, TCA as stop codon) and human genomic DNA as
template. TaqPlus Precision<sup>TM</sup> DNA polymerase (Stratagene) was used for the amplification
by the following cycle with step 2 to step 4 repeated 35 times: 94°C, 3 minutes; 94°C, 1
minute; 55°C, 1 minute; 72°C, 2 minutes; 72°C, 10 minutes.

A 1.3 Kb PCR fragment of predicated size was isolated and digested with Hind III 15 and Xba I, cloned into the pRC/CMV2 vector (Invitrogen) at the Hind III and Xba I sites and completely sequenced (SEQ.ID.NO.:1) using the T7 DNA Sequenase™ kit (Amsham).

#### j. hRUP3 (Seq. Id. Nos.:7 & 8)

The full length hRUP3 was cloned by PCR using the hRUP3 specific primers 5'-GTCCTGCCACTTCGAGACATGG-3' SEQ.ID.NO.:71 (sense, ATG as initiation codon) and 5'-20 GAAACTTCTCTGCCCTTACCGTC-3' SEQ.ID.NO.:72 (antisense, 3' of stop codon TAA) and human genomic DNA as template. TaqPlus Precision<sup>TM</sup> DNA polymerase (Stratagene) was used for the amplification by the following cycle with step 2 to step 4 repeated 35 times: 94°C, 3 minutes; 94°C, 1 minute; 58°C, 1 minute; 72°C, 2 minutes; 72°C, 10 minutes

A 1.0 Kb PCR fragment of predicated size was isolated and cloned into the pCRII-TOPOTM vector (Invitrogen) and completely sequenced (SEQ.ID.NO.: 7)using the T7 DNA sequenase kit (Amsham).

# Example 2 5 RECEPTOR EXPRESSION

Although a variety of cells are available to the art for the expression of proteins, it is most preferred that mammalian cells be utilized. The primary reason for this is predicated upon practicalities, *i.e.*, utilization of, *e.g.*, yeast cells for the expression of a GPCR, while possible, introduces into the protocol a non-mammalian cell which may not (indeed, in the 10 case of yeast, does not) include the receptor-coupling, genetic-mechanism and secretary pathways that have evolved for mammalian systems – thus, results obtained in non-mammalian cells, while of potential use, are not as preferred as that obtained from mammalian cells. Of the mammalian cells, COS-7, 293 and 293T cells are particularly preferred, although the specific mammalian cell utilized can be predicated upon the particular 15 needs of the artisan. The general procedure for expression of the disclosed GPCRs is as follows.

On day one, 1X10<sup>7</sup> 293T cells per 150mm plate were plated out. On day two, two reaction tubes will be prepared (the proportions to follow for each tube are per plate): tube A will be prepared by mixing 20µg DNA (e.g., pCMV vector; pCMV vector with receptor 20 cDNA, etc.) in 1.2ml serum free DMEM (Irvine Scientific, Irvine, CA); tube B will be prepared by mixing 120µl lipofectamine (Gibco BRL) in 1.2ml serum free DMEM. Tubes A and B are admixed by inversions (several times), followed by incubation at room temperature for 30-45min. The admixture can be referred to as the "transfection mixture". Plated 293T cells are washed with 1XPBS, followed by addition of 10ml serum free DMEM.

2.4ml of the transfection mixture will then be added to the cells, followed by incubation for 4hrs at 37°C/5% CO<sub>2</sub>. The transfection mixture was then be removed by aspiration, followed by the addition of 25ml of DMEM/10% Fetal Bovine Serum. Cells will then be incubated at 37°C/5% CO<sub>2</sub>. After 72hr incubation, cells can then be harvested and utilized for analysis.

#### 5 Example 3

#### TISSUE DISTRIBUTION OF THE DISCLOSED HUMAN GPCRS

Several approaches can be used for determination of the tissue distribution of the GPCRs disclosed herein.

#### 1. Dot-Blot Analysis

Using a commercially available human-tissue dot-blot format, endogenous orphan GPCRs were probed for a determination of the areas where such receptors are localized. cDNA fragments from the GPCRs of Example 1 (radiolabelled) were (or can be) used as the probe: radiolabeled probe was (or can be) generated using the complete receptor cDNA (excised from the vector) using a Prime-It II<sup>TM</sup> Random Primer Labeling Kit (Stratagene, 15 #300385), according to manufacturer's instructions. A human RNA Master Blot<sup>TM</sup> (Clontech, #7770-1) was hybridized with the endogenous human GPCR radiolabeled probe and washed under stringent conditions according manufacturer's instructions. The blot was exposed to Kodak BioMax<sup>TM</sup> Autoradiography film overnight at -80°C. Results are summarized for several receptors in Table B and C (see Figures 1A and 1B for a grid 20 identifying the various tissues and their locations, respectively). Exemplary dot-blots are provided in Figure 2A and 2B for results derived using hCHN3 and hCHN8, respectively.

#### TABLE B

ORPHAN GPCR

Tissue Distribution (highest levels, relative to other tissues in the dot-blot)

· · · · · · · · · · · · · · · · · · ·	hGPCR27	Fetal brain, Putamen, Pituitary gland, Caudate nucleus			
	hARE-1	Spleen, Peripheral leukocytes, Fetal spleen			
e e e e e e e e e e e e e e e e e e e	hPPR1	Pituitary gland, Heart, salivary gland, Small intestine, Testis			
	hRUP3	Pancreas			
5	hCHN3	Fetal brain, Putamen, Occipital cortex			
in the second se	hCHN9	Pancreas, Small intestine, Liver			
v en en	hCHN10	Kidney, Thryoid			

#### TABLE C

	Oi	RPHAN GPCR	Tissue Distribution (highest levels, relative to other tissues in the dot-blot)
10		hARE-3	Cerebellum left, Cerebellum right, Testis, Accumbens
		hGPCR3	Corpus collusum, Caudate nucleus, Liver, Heart, Inter- Ventricular Septum
		hARE-2	Cerebellum left, Cerebellum right, Substantia
12.	· · · · ·	hCHN8	Cerebellum left, Cerebellum right, Kidney, Lung

## 2. RT-PCR

## a. hRUP3

To ascertain the tissue distribution of hRUP3 mRNA, RT-PCR was performed using hRUP3-specific primers and human multiple tissue cDNA panels (MTC, Clontech) as templates. Taq DNA polymerase (Stratagene) was utilized for the PCR reaction, using the following reaction cycles in a 40ul reaction: 94°C for 2 min; 94°C for 15 sec; 55°C for 30 20 sec; 72°C for 1 min; 72° C, for 10 min. Primers were as follows:

- 5'-GACAGGTACCTTGCCATCAAG-3' (SEQ.ID.NO.: 61; sense)
- 5'-CTGCACAATGCCAGTGATAAGG-3' (SEQ.ID.NO.: 62; antisense).

20ul of the reaction was loaded onto a 1% agarose gel; results are set forth in Figure 3.

As is supported by the data of Figure 3, of the 16 human tissues in the cDNA panel utilized (brain, colon, heart, kidney, lung, ovary, pancreas, placenta, prostate, skeleton, small intestine, spleen, testis, thymus leukocyte, and liver) a single hRUP3 band is evident only from the pancreas. Additional comparative analysis of the protein sequence of hRUP3 with 5 other GPCRs suggest that hRUP3 is related to GPCRs having small molecule endogenous ligand such that it is predicted that the endogenous ligand for hRUP3 is a small molecule.

#### b. hRUP4

RT-PCR was performed using hRUP4 oligo's 8 and 4 as primers and the human multiple tissue cDNA panels (MTC, Clontech) as templates. Taq DNA polymerase 10 (Stratagene) was used for the amplification in a 40ul reaction by the following cycles: 94°C for 30 seconds, 94°C for 10 seconds, 55°C for 30 seconds, 72°C for 2 minutes, and 72°C for 5 minutes with cycles 2 through 4 repeated 30 times.

20 µl of the reaction were loaded on a 1% agarose gel to analyze the RT-PCR products, and hRUP4 mRNA was found expressed in many human tissues, with the strongest 15 expression in heart and kidney. (see, Figure 4). To confirm the authenticity of the PCR fragments, a 300 bp fragment derived from the 5' end of hRUP4 was used as a probe for the Southern Blot analysis. The probe was labeled with <sup>32</sup>P-dCTP using the Prime-It II<sup>™</sup>M Random Primer Labeling Kit (Stratagene) and purified using the ProbeQuant G-50 micro columns (Amersham). Hybridization was done overnight at 42° C following a 12 hr pre-20 hybridization. The blot was finally washed at 65°C with 0.1 x SSC. The Southern blot did confirm the PCR fragments as hRUP4.

#### c. hRUP5

RT-PCR was performed using the following hRUP5 specific primers:

- 5'-CTGACTTCTTGTTCCTGGCAGCAGCGG-3' (SEQ.ID.NO.: 63; sense)
- 5'-AGACCAGCCAGGGCACGCTGAAGAGTG-3' (SEQ.ID.NO.: 64; antisense)

and the human multiple tissue cDNA panels (MTC, Clontech) as templates. Taq DNA 5 polymerase (Stratagene) was used for the amplification in a 40ul reaction by the following cycles: 94°C for 30 sec, 94°C for 10 sec, 62°C for 1.5 min, 72°C for 5 min, and with cycles 2 through 3 repeated 30 times. 20 µl of the reaction were loaded on a 1.5% agarose gel to analyze the RT-PCR products, and hRUP5 mRNA was found expressed only in the peripheral blood leukocytes (*data not shown*).

#### d. hRUP6

RT-PCR was applied to confirm the expression and to determine the tissue distribution of hRUP6. Oligonucleotides used, based on an alignment of AC005871 and GPR66 segments, had the following sequences:

- 5'-CCAACACCAGCATCCATGGCATCAAG-3' (SEQ.ID.NO.: 73; sense),
- and the human multiple tissue cDNA panels (MTC, Clontech) were used as templates.

  PCR was performed using TaqPlus Precision™ polymerase (Stratagene; manufacturing instructions will be followed) in a 40ul reaction by the following cycles: 94°C for 30 sec; 94°C 5 sec; 66°C for 40 sec, 72°C for 2.5 min, and 72°C for 7 min. Cycles 2 through 4 20 were repeated 30 times.

20 ul of the reaction were loaded on a 1.2% agarose gel to analyze the RT-PCR products, and a specific 760bp DNA fragment representing hRUP6 was expressed predominantly in the thymus and with less expression in the heart, kidney, lung, prostate small intestine and testis. (see, Figure 5).

It is intended that each of the patents, applications, and printed publications mentioned in this patent document be hereby incorporated by reference in their entirety.

As those skilled in the art will appreciate, numerous changes and modifications may be made to the preferred embodiments of the invention without departing from the 5 spirit of the invention. It is intended that all such variations fall within the scope of the invention and the claims that follow.

Although a variety of Vectors are available to those in the art, for purposes of utilization for both endogenous and non-endogenous human GPCRs, it is most preferred that the Vector utilized be pCMV. This vector was deposited with the American Type 10 Culture Collection (ATCC) on October 13, 1998 (10801 University Blvd., Manassas, VA 20110-2209 USA) under the provisions of the Budapest Treaty for the International Recognition of the Deposit of Microorganisms for the Purpose of Patent Procedure. The DNA was tested by the ATCC and determined to be. The ATCC has assigned the following deposit number to pCMV: ATCC #203351.

#### **CLAIMS**

## What is claimed is:

- 1. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 1.
- 5 2. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 1 comprising SEQ.ID.NO.: 2.
  - 3. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:1.
  - 4. A Host Cell comprising the Plasmid of claim 3.
- 5. A cDNA encoding a human G protein-coupled receptor comprising 10 SEQ.ID.NO.: 3.
  - 6. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 3 comprising SEQ.ID.NO.: 4.
    - 7. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:3.
    - 8. A Host Cell comprising the Plasmid of claim 7.
- 9. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 5.
  - 10. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 5 comprising SEQ.ID.NO.: 6.
    - 11. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:5.
- 20 12. A Host Cell comprising the Plasmid of claim 11.
  - 13. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 7.

- 14. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 7 comprising SEQ.ID.NO.: 8.
  - 15. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:7.
  - 16. A Host Cell comprising the Plasmid of claim 15.
- 17. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 9.
- 18. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 9 comprising SEQ.ID.NO.: 10.
  - 19. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:9.
- 10 20. A Host Cell comprising the Plasmid of claim 19.
  - 21. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 11.
  - 22. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 11 comprising SEQ.ID.NO.:12.
- 15 23. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:11.
  - 24. A Host Cell comprising the Plasmid of claim 23.
  - 25. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 13.
- 26. A human G protein-coupled receptor encoded by the cDNA of 20 SEQ.ID.NO.: 13 comprising SEQ.ID.NO.: 14.
  - 27. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:13.
  - 28. A Host Cell comprising the Plasmid of claim 27.
  - 29. A cDNA encoding a human G protein-coupled receptor comprising

## SEQ.ID.NO.: 15.

- 30. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 15 comprising SEQ.ID.NO.: 16.
  - 31. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:15.
- 5 32. A Host Cell comprising the Plasmid of claim 31.
  - 33. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 17.
  - 34. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 17 comprising SEQ.ID.NO.: 18.
- 10 35. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:17.
  - 36. A Host Cell comprising the Plasmid of claim 35.
  - 37. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 19.
- 38. A human G protein-coupled receptor encoded by the cDNA of 15 SEQ.ID.NO.: 19 comprising SEQ.ID.NO.: 20.
  - 39. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:19.
  - 40. A Host Cell comprising the Plasmid of claim 39.
  - 41. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 21.
- 20 42. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 21 comprising SEQ.ID.NO.: 22.
  - 43. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:21.
  - 44. A Host Cell comprising the Plasmid of claim 43.

- 45. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 23.
- 46. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 23 comprising SEQ.ID.NO.: 24.
- 5 47. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.: 23.
  - 48. A Host Cell comprising the Plasmid of claim 47.
  - 49. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 25.
- 50. A human G protein-coupled receptor encoded by the cDNA of 10 SEQ.ID.NO.: 25 comprising SEQ.ID.NO.: 26.
  - 51. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:25.
  - 52. A Host Cell comprising the Plasmid of claim 51.
  - 53. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 27.
- 15 54. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 27 comprising SEQ.ID.NO.: 28.
  - 55. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:27.
  - 56. A Host Cell comprising the Plasmid of claim 55.
- 57. A cDNA encoding a human G protein-coupled receptor comprising 20 SEQ.ID.NO.: 29.
  - 58. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 29 comprising SEQ.ID.NO.: 30.
    - 59. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:29.

- 60. A Host Cell comprising the Plasmid of claim 59.
- 61. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 31.
- 62. A human G protein-coupled receptor encoded by the cDNA of 5 SEQ.ID.NO.: 31 comprising SEQ.ID.NO.: 32.
  - 63. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:31.
  - 64. A Host Cell comprising the Plasmid of claim 63.
- 65. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 33.
- 10 66. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 33 comprising SEQ.ID.NO.: 34.
  - 67. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:33.
  - 68. A Host Cell comprising the Plasmid of claim 67.
- 69. A cDNA encoding a human G protein-coupled receptor comprising 15 SEQ.ID.NO.: 35.
  - 70. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 35 comprising SEQ.ID.NO.: 36.
    - 71. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:35.
    - 72. A Host Cell comprising the Plasmid of claim 71.
- 73. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 37.
  - 74. A human G protein-coupled receptor encoded by the cDNA.of SEQ.ID.NO.: 37 comprising SEQ.ID.NO.: 38.

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- 36 -

75. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:37.

76. A Host Cell comprising the Plasmid of claim 75.

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O	Me		mpG TOTO	*	Stomach	,-	Mannary	Gland	Bone	Marrow				
7	Hippocampus	T T T	Spinal Cord		Prostate		Salivary	Gland	Lymph	Node		Fetal	Time	Q.m.
9	Frontal	Cortex	Accumbens		Uterus		Thyroid		Peripheral	Leukocyte		Fetal	Thymns	
5	Cerabral	Cortex	Thalamus		Bladder	1 =	Adrenal	Gland	Thymus			Fetal	Spleen	
4	Cerebellum		Temporal	Cortex	Colon	)	Pituitary		Spleen		Placenta	Fetal	Liver	
3	Caudate	Nucleus	Substantia	Nigra	Skeletal	Muscle	Pancreas	*	Small	Intestine	Trachea	Fetal	Kidney	
2	Amygdala		Putamen	*	Aorta		Ovary		Liver		Lung	Fetal	Heart	
-			Occipital	Cortex	Heart		Testis	-	Kidney		Appendix	Fetal	Brain	
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	=	Fetal	Brain	Fotol	3 -	1631	– ਮੁਹਤ	Kidincy	Fetal	Live	Fetal	Spleen		Fetal	Thymus		Fetal	Ling	0					
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	6	Liver		Pancreas	· 2	Adrenal	ק - ספר ל	Cland	Ihyroid		Salivary	Cland		Mammary	Gland									
	8	Lung		Placenta		Bladder		1 1	Cterus		Prostate			Testis			Ovary							- <u></u>
	7	Kidney		Skeletal	Muscle	Spleen		F	ınymus		Peripheral	Leukocyte		Lymph	Node		Bone	Marrow	**	4 22	Traches	4		
	9	Colon	Iransverse	Colon	Descriding	Rectum									*									
	<b>.</b>	Esophagus		Stomach		Duodenum		Teinmim		-	TIGETH 1			Ilocecum			Appendix			, .	Colon	Ascending	0	
,	4	Heart		Aorta	-)(-	Atrium	Left	Afrim	Right	Vication	venucie	Tell Tell		Ventricle	Kight		Inter	Ventricular	Septum		Apex of	the Heart	6	
2		Substantia	n Andra	Accumbens	*	Thalamus		Pituitary	Gland	Chinal		<b>g</b>											0.	
2	1	Cerebellum	1	Carecellum	Right	Corpus	Callosum	Amygdala		Cardete	Niclers	TOPPICATION	11	Hippocampus		11.	Ivadulla	CONTINUE			Putamen			
			Construct	A William	Cortex	Frontal	Cortex	Parietal	Lobe	Occipital	Cortex			Cortex	<b>3</b>	Paracentur	Je maranan	Cyrick of	E CODE	Cortex	Pons			

FIG. 108

3/4

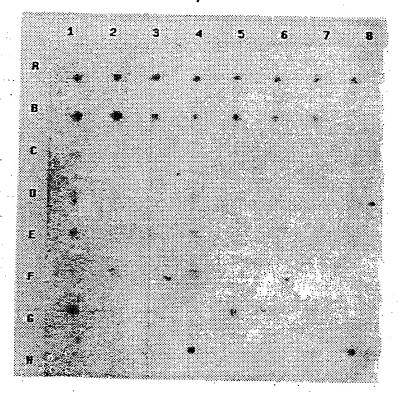


FIG. 2A

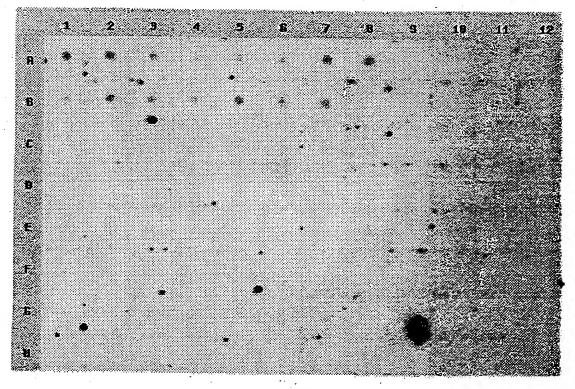


FIG. 2B

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Thymus
Testis
Spleen
Small Intestine
Skeletal Muscle
Prostate

Placenta

Pancreas 40vary

lung Liver Leukocyte

Kidney

Heart Colon

Brain

PCT/US99/23687

5

#### SEQUENCE LISTING

#### (1) GENERAL INFORMATION:

(i) APPLICANT: Chen, Ruoping
Dang, Huong T.
Liaw, Chen W.
Lin, I-Lin

- (ii) TITLE OF INVENTION: Human Orphan G Protein-Coupled Receptors
- (iii) NUMBER OF SEQUENCES: 74
- 10 (iv) CORRESPONDENCE ADDRESS:
  - (A) ADDRESSEE: Arena Pharmaceuticals, Inc.
  - (B) STREET: 6166 Nancy Ridge Drive
  - (C) CITY: San Diego
  - (D) STATE: CA
- 15 (E) COUNTRY: USA
  - (F) ZIP: 92121
  - (v) COMPUTER READABLE FORM:
    - (A) MEDIUM TYPE: Floppy disk
    - (B) COMPUTER: IBM PC compatible
- 20 (C) OPERATING SYSTEM: PC-DOS/MS-DOS
  - (D) SOFTWARE: PatentIn Release #1.0, Version #1.30
  - (vi) CURRENT APPLICATION DATA:
    - (A) APPLICATION NUMBER: US
    - (B) FILING DATE:
- 25 (C) CLASSIFICATION:
  - (viii) ATTORNEY/AGENT INFORMATION:
    - (A) NAME: Burgoon, Richard P.
      - (B) REGISTRATION NUMBER: 34,787
    - (ix) TELECOMMUNICATION INFORMATION:

(A) TELEPHONE: (858)453-7200

- (B) TELEFAX: (858)453-7210
- (2) INFORMATION FOR SEQ ID NO:1:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 1260 base pairs
- 35 (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
  - (ii) MOLECULE TYPE: DNA (genomic)
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

GTGTATGAAA ACACCTACAT GAATATTACA CTCCCTCCAC CATTCCAGCA TCCTGACCTC 120 AGTCCATTGC TTAGATATAG TTTTGAAACC ATGGCTCCCA CTGGTTTGAG TTCCTTGACC 180 GTGAATAGTA CAGCTGTGCC CACAACACCA GCAGCATTTA AGAGCCTAAA CTTGCCTCTT 240 CAGATCACCC TTTCTGCTAT AATGATATTC ATTCTGTTTG TGTCTTTTCT TGGGAACTTG 300 5 GTTGTTTGCC TCATGGTTTA CCAAAAAGCT GCCATGAGGT CTGCAATTAA CATCCTCCTT 360 GCCAGCCTAG CTTTTGCAGA CATGTTGCTT GCAGTGCTGA ACATGCCCTT TGCCCTGGTA 420 ACTATTCTTA CTACCCGATG GATTTTTGGG AAATTCTTCT GTAGGGTATC TGCTATGTTT 480 TTCTGGTTAT TTGTGATAGA AGGAGTAGCC ATCCTGCTCA TCATTAGCAT AGATAGGTTC 540 CTTATTATAG TCCAGAGGCA GGATAAGCTA AACCCATATA GAGCTAAGGT TCTGATTGCA 600 10 GTTTCTTGGG CAACTTCCTT TTGTGTAGCT TTTCCTTTAG CCGTAGGAAA CCCCGACCTG 660 CAGATACCTT CCCGAGCTCC CCAGTGTGTG TTTGGGTACA CAACCAATCC AGGCTACCAG 720 GCTTATGTGA TTTTGATTTC TCTCATTTCT TTCTTCATAC CCTTCCTGGT AATACTGTAC 780 TCATTTATGG GCATACTCAA CACCCTTCGG CACAATGCCT TGAGGATCCA TAGCTACCCT 840 GAAGGTATAT GCCTCAGCCA GGCCAGCAAA CTGGGTCTCA TGAGTCTGCA GAGACCTTTC 900 15 CAGATGAGCA TTGACATGGG CTTTAAAACA CGTGCCTTCA CCACTATTTT GATTCTCTTT 960 GCTGTCTTCA TTGTCTGCTG GGCCCCATTC ACCACTTACA GCCTTGTGGC AACATTCAGT1020

AAGCACTTT ACTATCAGCA CAACTTTTT GAGATTAGCA CCTGGCTACT GTGGCTCTGC1080
TACCTCAAGT CTGCATTGAA TCCGCTGATC TACTACTGGA GGATTAAGAA ATTCCATGAT1140
20 GCTTGCCTGG ACATGATGCC TAAGTCCTTC AAGTTTTTGC CGCAGCTCCC TGGTCACACA1200
AAGCGACGGA TACGTCCTAG TGCTGTCTAT GTGTGGGG AACATCGGAC GGTGGTGGA1260

- (3) INFORMATION FOR SEQ ID NO:2:
  - (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 419 amino acids
  - (B) TYPE: amino acid
    - (C) STRANDEDNESS: single
    - (D) TOPOLOGY: linear
    - (ii) MOLECULE TYPE: DNA (genomic)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:
- Met Val Phe Ser Ala Val Leu Thr Ala Phe His Thr Gly Thr Ser Asn

		Thr	Thi	r Phe	20	l Va]	L Ty:	r Glı	ı Asr	1 Thi 25	Tyr	Met	Ası	ı Ile	Thi 30	Leu	Pro
		Pro	Pro	Phe 35	Glr	ı His	Pro	) Asp	Let 40	Sei	Pro	Let	Let	1 Arc	Туг	Ser	Phe
5		Glu	Thr	. Met	: Ala	Pro	Thr		Leu	Ser	Ser	Lev		,	. Asr	. Ser	Thr
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10		Gln	Ile	Thr	Leu	Ser 85	Ala	lle	Met	Ile	Phe 90	Ile	Leu	Phe	Val	Ser 95	Phe
		Leu	Gly	Asn	Leu 100	Val	Val	Cys	Leu	Met	Val	Tyr	Gln	Lys	Ala		Met
	ē .	Arg	Ser	Ala 115	Ile	Asn	Ile	Leu	Leu 120		Ser	Leu	Ala	Phe 125	Ala	Asp	Met
15		Leu	Leu 130	Ala	Val	Leu	Asn	Met 135	Pro	Phe	Ala	Leu	Val		Ile	Leu	Thr
		Thr 145	Arg	Trp	Ile	Phe	Gly 150		Phe	Phe	Cys	Arg 155	Val	Ser	Ala	Met	Phe
20		Phe	Trp	Leu	Phe	Val 165	Ile	Glu	Gly	Val	Ala 170	Ile	Leu	Leu	Ile	Ile 175	Ser
		Ile	Asp	Arg	Phe 180	Leu	Ile	Ile	Val	Gln 185	Arg	Gln	Asp	Lys	Leu 190	Asn	Pro
		Tyr	Arg	Ala 195	Lys	Val	Leu	Ile	Ala 200	Val	Ser.	Trp	Ala	Thr 205	Ser	Phe	Сув
25		Val	Ala 210	Phe	Pro	Leu .,	Ala	Val 215	Gly	Asn	Pro	Asp	Leu 220		Ile	Pro	Ser
*	•	Arg 225	Ala	Pro	Gln	Cys	Val 230	Phe	Gly	Tyr	Thr	Thr 235	Aşn	Pro	Gly	Tyr	Gln 240
3Ô		Ala	Tyr	Val	Ile	Leu 245	Ile	Ser	Leu	Ile	Ser 250	Phe	Phe	Ile	Pro	Phe 255	Leu
-		Val	Ile	Leu	Tyr 260	Ser	Phe	Met	Gly	Ile 265	Leu	Asn	Thr	Leu	Arg 270	His	Asn
		Ala	Leu	Arg 275	Ile	His	Ser	Tyr	Pro 280	Glu	Gly	Ile	Суѕ	Leu 285	Ser	Gln	Ala
35		Ser	Lys 290	Lëu	Gly	Leu	Met	Ser 295	Leu	Gln	Arg	Pro	Phe	Gln	Met	Ser	Ile
٠.		<i>l</i> .sp	Met	Gly	Phe	Lys	Thr	Arg	Ala	Phe	Thr	Thr	Ile	Leu	Ile	Leu	Phe

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Ala Val Phe Ile Val Cys Trp Ala Pro Phe Thr Thr Tyr Ser Leu Val
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Ala Thr Phe Ser Lys His Phe Tyr Tyr Gln His Asn Phe Phe Glu Ile
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Ser Thr Trp Leu Leu Trp Leu Cys Tyr Leu Lys Ser Ala Leu Asn Pro
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Leu Ile Tyr Tyr Trp Arg Ile Lys Lys Phe His Asp Ala Cys Leu Asp
370
375
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Met Met Pro Lys Ser Phe Lys Phe Leu Pro Gln Leu Pro Gly His Thr
385
390
395
400

Lys Arg Arg Ile Arg Pro Ser Ala Val Tyr Val Cys Gly Glu His Arg

Thr Val Val

15 -

# (4) INFORMATION FOR SEQ ID NO:3:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1119 base pairs
  - (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
  - (ii) MOLECULE TYPE: DNA (genomic)
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

ATGTTAGCCA ACAGCTCCTC AACCAACAGT TCTGTTCTCC CGTGTCCTGA CTACCGACCT 60

25 ACCCACCGCC TGCACTTGGT GGTCTACAGC TTGGTGCTGG CTGCCGGGGCT CCCCCTCAAC 120

GCGCTAGCCC TCTGGGTCTT CCTGCGCGCG CTGCGCGTGC ACTCGGTGGT GAGCGTGTAC 180

ATGTGTAACC TGGCGGCCAG CGACCTGCTC TTCACCCTCT CGCTGCCCGT TCGTCTCTCC 240

TACTACGCAC TGCACCACTG GCCCTTCCCC GACCTCCTGT GCCAGACGAC GGGCGCCATC 300

TTCCAGATGA ACATGTACGG CAGCTGCATC TTCCTGATGC TCATCAACGT GGACCGCTAC 360

30 GCCGCCATCG TGCACCCGCT GCGACTGCGC CACCTGCGGC GGCCCGCGT GGCGCGGCTG 420

CTCTGCCTGG GCGTGTGGGC GCTCATCCTG GTGTTTGCCG TGCCCGCCGC CCGCGTGCAC 480

AGGCCCTCGC GTTGCCGCTA CCGGGACCTC GAGGTGCGCC TATGCTTCGA GAGCTTCAGC 540

GACGAGCTGT GGAAAGGCAG GCTGCTGCCC CTCGTGCTGC TGGCCGAGGC GCTGGGCTTC 600

CTGCTGCCCC TGGCGGCGGT GGTCTACTCG TCGGGCCGAG TCTTCTGGAC GCTGGCGCGC 660

CCCGACGCCA CGCAGAGCCA GCGGCGGCG AAGACCGTGC GCCTCCTGCT GGCTAACCTC 720

GTCATCTTCC TGCTGTGCTT CGTGCCCTAC AACAGCACGC TGGCGGTCTA CGGGCTGCTG 780

CGGAGCAAGC TGGTGGCGG CAGCGTGCCT GCCCGCGATC GCGTGCGCGG GGTGCTGATG 840

5 GTGATGGTGC TGCTGGCCGG CGCCAACTGC GTGCTGGACC CGCTGGTGTA CTACTTTAGC 900

GCCGAGGGCT TCCGCAACAC CCTGCGCGGC CTGGGCACTC CGCACCGGGC CAGGACCTCG 960

GCCACCAACG GGACGCGGG GGCGCTCGCG CAATCCGAAA GGTCCGCCGT CACCACCGAC1020

GCCACCAGGC CGGATGCCGC CAGTCAGGGG CTGCTCCGAC CCTCCGACTC CCACTCTCTG1080

TCTTCCTTCA CACAGTGTCC CCAGGATTCC GCCCTCTGA

## 10 (5) INFORMATION FOR SEQ ID NO:4:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 372 amino acids
    - (B) TYPE: amino acid
    - (C) STRANDEDNESS:
- 15 (D) TOPOLOGY: not relevant
  - (ii) MOLECULE TYPE: protein
    - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

Met Leu Ala Asn Ser Ser Ser Thr Asn Ser Ser Val Leu Pro Cys Pro-

- 20 Asp Tyr Arg Pro Thr His Arg Leu His Leu Val Val Tyr Ser Leu Val
  20 25 30
  - Leu Ala Ala Gly Leu Pro Leu Asn Ala Leu Ala Leu Trp Val Phe Leu 35 40 45
- Arg Ala Leu Arg Val His Ser Val Val Ser Val Tyr Met Cys Asn Leu
  50 55 60
  - Ala Ala Ser Asp Leu Leu Phe Thr Leu Ser Leu Pro Val Arg Leu Ser 65 70 75 80
  - Tyr Tyr Ala Leu His His Trp Pro Phe Pro Asp Leu Leu Cys Gln Thr
    85 90 95
- Thr Gly Ala Ile Phe Gln Met Asn Met Tyr Gly Ser Cys Ile Phe Leu
  100 105 110
  - Met Leu Ile Asn Val Asp Arg Tyr Ala Ala Ile Val His Pro Leu Arg

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			*.	,		4.1.										
	Leu	Arg	His	Leu	Arg	Arg	Pro	Arg	Val	Ala	Arg	Leu	Leu	Cys	Leu	Gly
5	*	130					135					140		- 7		
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	Val	Trp	Ala	Leu	Ile	Leu	Val	Phe	Ala	Val	Pro	Ala	Ala	Arg	Val	His
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	Glu	Ser	Dhe	Ser	Δen	Glin	T.A11	Trn	Tye	Gly	7~~	T 011	Ton	Dwo	Leu	Wal.
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		Ser	Gln	Arg	Arg	Arg	Lys	Thr	Val	Arg	Leu	Leu	Leu	Ala	Asn	Leu
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	Asp	Arg	Val	Arg	Gly	Val	Leu	Met	Val	Met	Val	Leu	Leu	Ala	Gly	Ala
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	Asn		Val	Leu	Asp	Pro		Val	Tyr	Tyr	Phe		Ala	Glu	Gly	Phe
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	Val	Thr	Thr	Asp	Ala	Thr	Arg	Pro	Asp	Ala	Ala	Ser	Gln	Gly	Leu	Leu
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	Arg	Pro		Asp	Ser	His	Ser		Ser	Ser	Phe	Thr.		Cys	Pro	Gln
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30	Asp	Ser		Leu						0						•
30	Asp	Ser 370		Leu	*					0						*

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1107 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: single

## (ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

ATGGCCAACT CCACAGGGCT GAACGCCTCA GAAGTCGCAG GCTCGTTGGG GTTGATCCTG 60 GCAGCTGTCG TGGAGGTGGG GGCACTGCTG GGCAACGGCG CGCTGCTGGT CGTGGTGCTG 120 5 CGCACGCCGG GACTGCGCGA CGCGCTCTAC CTGGCGCACC TGTGCGTCGT GGACCTGCTG 180 GCGGCCGCCT CCATCATGCC GCTGGGCCTG CTGGCCGCAC CGCCGCCCGG GCTGGGCCGC 240 GTGCGCCTGG GCCCGCGCC ATGCCGCGCC GCTCGCTTCC TCTCCGCCGC TCTGCTGCCG 300 GCCTGCACGC TCGGGGTGGC CGCACTTGGC CTGGCACGCT ACCGCCTCAT CGTGCACCCG 360 CTGCGGCCAG GCTCGCGGCC GCCGCCTGTG CTCGTGCTCA CCGCCGTGTG GGCCGCGGCG 420 10 GGACTGCTGG GCGCGCTCTC CCTGCTCGGC CCGCCGCCCG CACCGCCCCC TGCTCCTGCT 480 CGCTGCTCGG TCCTGGCTGG GGGCCTCGGG CCCTTCCGGC CGCTCTGGGC CCTGCTGGCC 540 TTCGCGCTGC CCGCCCTCCT GCTGCTCGGC GCCTACGGCG GCATCTTCGT GGTGGCGCGT 600 CGCGCTGCCC TGAGGCCCCC ACGGCCGGCG CGCGGGTCCC GACTCCGCTC GGACTCTCTG 660 GATAGCCGCC TTTCCATCTT GCCGCCGCTC CGGCCTCGCC TGCCCGGGGG CAAGGCGGCC 720 15 CTGGCCCCAG CGCTGGCCGT GGGCCAATTT GCAGCCTGCT GGCTGCCTTA TGGCTGCGCG 780 TGCCTGGCGC CCGCAGCGCG GGCCGCGGAA GCCGAAGCGG CTGTCACCTG GGTCGCCTAC 840 TCGGCCTTCG CGGCTCACCC CTTCCTGTAC GGGCTGCTGC AGCGCCCCGT GCGCTTGGCA 900 CTGGGCCGCC TCTCTCGCCG TGCACTGCCT GGACCTGTGC GGGCCTGCAC TCCGCAAGCC 960 TGGCACCCGC GGGCACTCTT GCAATGCCTC CAGAGACCCC CAGAGGGCCC TGCCGTAGGC1020 20 CCTTCTGAGG CTCCAGAACA GACCCCCGAG TTGGCAGGAG, GGCGGAGCCC CGCATACCAG1080 GGGCCACCTG AGAGTTCTCT CTCCTGA 1107

#### (7) INFORMATION FOR SEQ ID NO:6:

25

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 368 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS:
  - (D) TOPOLOGY: not relevant
- (ii) MOLECULE TYPE: protein
- (x1) SEQUENCE DESCRIPTION: SEQ ID NO:6:

*	Me 1	t Al	a As	n Se	r Thi	Gl:	y Lei	u Ası	n Ala	a Se:	r Gli	u Va	l Al	a Gly	Ser ::15	Leu
	Gl	y Le	u Il	e Lei 20	ı Ala	ı Ala	a Va	l Val	l Glu 25	ı Va	l Gly	γ Ala	a Lei	ı Leu 30	Gly	/ Asn
.5	Gl	y Al	a Le	u Let	, ,	. Va]	l Va]	L Let 40	ı Arg	Thi	r Pro	o Gly	/ Let 45	ı Arg	Asp	Ala
1		- 30					55					60	*	Ala	*	
10	. 03					70				*	75			Leu	·. / :	80
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	٠. `	130	·				135					140		Leu		
20	143		-		*	150				;	155		* : 10	Ala		160
			* 1		165					170					175	
25			i * *	180	· · · · · ·	٠	· · · · ·		185	, j				190 Pro		
		Ala	732			Arg	Leu	200					205	Ser		
30		210		• • •	Pro	Leu	215				Pro	220		Lys		
	-	Ala	Pro	Ala		230 Ala	Val	Gly			235 Ala	Ala	Суз	Trp 1	Leu	240 Pro
	Tyr	Gly	Cys			Leu	Ala	Pro		250 Ala	Arg	Ala		Glu <i>I</i>	255 Ala (	Glu
35	Ala	Ala	Val 275		Trp	Val	Ala			-	Phe .			270 His I	Pro 1	Phe
	Leu	Tyr	Gly	Leu i	Leu (	3ln	Arg	Pro	Val	Arg	Leu .			Gly A	\rg	Leu

290 295 300

Ser Arg Arg Ala Leu Pro Gly Pro Val Arg Ala Cys Thr Pro Gln Ala 305 310 315 320

Trp His Pro Arg Ala Leu Leu Gln Cys Leu Gln Arg Pro Pro Glu Gly
325 330 335

Pro Ala Val Gly Pro Ser Glu Ala Pro Glu Gln Thr Pro Glu Leu Ala 340 345 350

Gly Gly Arg Ser Pro Ala Tyr Gln Gly Pro Pro Glu Ser Ser Leu Ser 355 360 365

## 10 (8) INFORMATION FOR SEQ ID NO:7:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1008 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
- 5 (D) TOPOLOGY: linear
  - (ii) MOLECULE TYPE: DNA (genomic)

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

ATGGAATCAT CTTTCTCATT TGGAGTGATC CTTGCTGTCC TGGCCTCCT CATCATTGCT 60

ACTAACACAC TAGTGGCTGT GGCTGTGCTG CTGTTGATCC ACAAGAATGA TGGTGTCAGT 120

20 CTCTGCTTCA CCTTGAATCT GGCTGTGGCT GACACCTTGA TTGGTGTGGC CATCTCTGGC 180

CTACTCACAG ACCAGCTCTC CAGCCCTTCT CGGCCCACAC AGAAGACCCT GTGCAGCCTG 240

CGGATGGCAT TTGTCACTTC CTCCGCAGCT GCCTCTGTCC TCACGGTCAT GCTGATCACC 300

TTTGACAGGT ACCTTGCCAT CAAGCAGCCC TTCCGCTACT TGAAGATCAT GAGTGGGTTC 360

GTGGCCGGGG CCTGCATTGC CGGGCTGTGG TTAGTGTCTT ACCTCATTGG CTTCCTCCCA 420

25 CTCGGAATCC CCATGTTCCA GCAGACTGCC TACAAAGGGC AGTGCAGCTT CTTTGCTGTA 480

TTTTCACCCTC ACTTCGTGCT GACCCTCTCC TGCGTTGGCT TCTTCCCAGC CATGCTCCTC 540

AAGATGGAAC ATGCAGGAG CATGCTCAAG ATTGCCTCCA TGCACAGCCA GCAGATTCGA 600

AAGATGGAAC ATGCAGGAG CATGGCTGGA GGTTATCGAT CCCCACGGAC TCCCAGCGAC 660

TTCAAAAGCTC TCCGTACTGT GTCTGTTCTC ATTGGGAGCT TTGCTCTATC CTGGACCCCC 720

30 TTCCTTATCA CTGGCATTGT GCAGGTGGCC TGCCAGGAGT GTCACCCCCT CATCAGTGCT 780

GAACGGTACC TGTGGCTGCT CGGCGTGGGC AACTCCCTGC TCAACCCACT CATCTATGCC 840

TATTGGCAGA AGGAGGTGCG ACTGCAGCTC TACCACATGG CCCTAGGAGT GAAGAAGGTG 900
CTCACCTCAT TCCTCCTCTT TCTCTCGGCC AGGAATTGTG GCCCAGAGAG GCCCAGGGAA 960
AGTTCCTGTC ACATCGTCAC TATCTCCAGC TCAGAGTTTG ATGGCTAA 1008

- (9) INFORMATION FOR SEQ ID NO:8:
- 5 (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 335 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS:
  - (D) TOPOLOGY: not relevant
- 10 (ii) MOLECULE TYPE: protein
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:
  - Met Glu Ser Ser Phe Ser Phe Gly Val Ile Leu Ala Val Leu Ala Ser 1 10 15
- Leu Ile Ile Ala Thr Asn Thr Leu Val Ala Val Ala Val Leu Leu Leu 20 25 30
  - Ile His Lys Asn Asp Gly Val Ser Leu Cys Phe Thr Leu Asn Leu Ala
    35 40 45
  - Val Ala Asp Thr Leu Ile Gly Val Ala Ile Ser Gly Leu Leu Thr Asp 50 55 60
- 20 Gln Leu Ser Ser Pro Ser Arg Pro Thr Gln Lys Thr Leu Cys Ser Leu
  65 70 75 80
  - Arg Met Ala Phe Val Thr Ser Ser Ala Ala Ala Ser Val Leu Thr Val 85 90 95
- Met Leu Ile Thr Phe Asp Arg Tyr Leu Ala Ile Lys Gln Pro Phe Arg 100 105 110
  - Tyr Leu Lys Ile Met Ser Gly Phe Val Ala Gly Ala Cys Ile Ala Gly
    115 120 125
  - Leu Trp Leu Val Ser Tyr Leu Ile Gly Phe Leu Pro Leu Gly Ile Pro 130 135 140
- Met Phe Gln Gln Thr Ala Tyr Lys Gly Gln Cys Ser Phe Phe Ala Val 145 150 155 160
  - Phe His Pro His Phe Val Leu Thr Leu Ser Cys Val Gly Phe Phe Pro 165 170 175
- Ala Met Leu Leu Phe Val Phe Phe Tyr Cys Asp Met Leu Lys Ile Ala
  180 185 190

Ser Met His Ser Gln Gln Ile Arg Lys Met Glu His Ala Gly Ala Met 200 Ala Gly Gly Tyr Arg Ser Pro Arg Thr Pro Ser Asp Phe Lys Ala Leu 215 - 220 Arg Thr Val Ser Val Leu Ile Gly Ser Phe Ala Leu Ser Trp Thr Pro 230 Phe Leu Ile Thr Gly Ile Val Gln Val Ala Cys Gln Glu Cys His Leu 245 Tyr Leu Val Leu Glu Arg Tyr Leu Trp Leu Leu Gly Val Gly Asn Ser 10 265 Leu Leu Asn Pro Leu Ile Tyr Ala Tyr Trp Gln Lys Glu Val Arg Leu 280 Gln Leu Tyr His Met Ala Leu Gly Val Lys Lys Val Leu Thr Ser Phe Leu Leu Phe Leu Ser Ala Arg Asn Cys Gly Pro Glu Arg Pro Arg Glu 310 315 Ser Ser Cys His Ile Val Thr Ile Ser Ser Ser Glu Phe Asp Gly

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- (10) INFORMATION FOR SEQ ID NO:9:
- 20 (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1413 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- 25 (ii) MOLECULE TYPE: DNA (genomic)
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

ATGGACACTA CCATGGAAGC TGACCTGGGT GCCACTGGCC ACAGGCCCCG CACAGAGCTT 60
GATGATGAGG ACTCCTACCC CCAAGGTGGC TGGGACACGG TCTTCCTGGT GGCCCTGCTG 120
CTCCTTGGGC TGCCAGCCAA TGGGTTGATG GCGTGGCTGG CCGGCTCCCA GGCCCGGCAT 180
30 GGAGCTGGCA CGCGTCTGGC GCTGCTCCTG CTCAGCCTGG CCCTCTCTGA CTTCTTGTTC 240
CTGGCAGCAG CGGCCTTCCA GATCCTAGAG ATCCGGCATG GGGGACACTG GCCGCTGGGG 300
ACAGCTGCCT GCCGCTTCTA CTACTTCCTA TGGGGCGTGT CCTACTCCTC CGGCCTCTTC 360
CTGCTGGCCG CCCTCAGCCT CGACCGCTGC CTGCTGGCGC TGTGCCCACA CTGGTACCCT 420
GGGCACCGCC CAGTCCGCC GCCCTCTGG GTCTGCGCC GTGTCTGGGT GCTGGCCACA 480

CTCTTCAGCG TGCCCTGGCT GGTCTTCCCC GAGGCTGCCG TCTGGTGGTA CGACCTGGTC 540 ATCTGCCTGG ACTTCTGGGA CAGCGAGGAG CTGTCGCTGA GGATGCTGGA GGTCCTGGGG 600 GGCTTCCTGC CTTTCCTCCT GCTGCTCGTC TGCCACGTGC TCACCCAGGC CACAGCCTGT 660 CGCACCTGCC ACCGCCAACA GCAGCCCGCA GCCTGCCGGG GCTTCGCCCG TGTGGCCAGG 720 5 ACCATTCTGT CAGCCTATGT GGTCCTGAGG CTGCCCTACC AGCTGGCCCA GCTGCTCTAC 780 CTGGCCTTCC TGTGGGACGT CTACTCTGGC TACCTGCTCT GGGAGGCCCT GGTCTACTCC 840 GACTACCTGA TCCTACTCAA CAGCTGCCTC AGCCCCTTCC TCTGCCTCAT GGCCAGTGCC 900 GACCTCCGGA CCCTGCTGCG CTCCGTGCTC TCGTCCTTCG CGGCAGCTCT CTGCGAGGAG 960 CGGCCGGGCA GCTTCACGCC CACTGAGCCA CAGACCCAGC TAGATTCTGA GGGTCCAACT1020 10 CTGCCAGAGC CGATGGCAGA GGCCCAGTCA CAGATGGATC CTGTGGCCCA GCCTCAGGTG1080 AACCCCACAC TCCAGCCACG ATCGGATCCC ACAGCTCAGC CACAGCTGAA CCCTACGGCC1140 CAGCCACAGT CGGATCCCAC AGCCCAGCCA CAGCTGAACC TCATGGCCCA GCCACAGTCA1200 GATTCTGTGG CCCAGCCACA GGCAGACACT AACGTCCAGA CCCCTGCACC TGCTGCCAGT1260 TCTGTGCCCA GTCCCTGTGA TGAAGCTTCC CCAACCCCAT CCTCGCATCC TACCCCAGGG1320 15 GCCCTTGAGG ACCCAGCCAC ACCTCCTGCC TCTGAAGGAG AAAGCCCCAG CAGCACCCCG1380 CCAGAGGCGG CCCCGGGCGC AGGCCCCACG TGA 1413

## (11) INFORMATION FOR SEQ ID NO:10:

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- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 468 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS:
  - (D) TOPOLOGY: not relevant
- (ii) MOLECULE TYPE: protein
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:
- Met Asp Thr Thr Met Glu Ala Asp Leu Gly Ala Thr Gly His Arg Pro

  1 5 10 15
  - Arg Thr Glu Leu Asp Asp Glu Asp Ser Tyr Pro Gln Gly Gly Trp Asp
    20 25 30
- Thr Val Phe Leu Val Ala Leu Leu Leu Gly Leu Pro Ala Asn Gly
  30 35 40 45

Leu Met Ala Trp Leu Ala Gly Ser Gln Ala Arg His Gly Ala Gly Thr

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0		Arg	, Lei	ı Ala	a Leu	Leu	Lei	ı Let	ı Sez	Lei	u Ala	Lei	ı Sei	r Asp	) Phe	Lei	ı Pho
		65					70					75			, .		80
٠,		Leu	i Ala	. Ala	Ala	Ala	Phe	Gln	Ile	Let	ı Gla	1 714	 • Arc	, His		G1.	, ui,
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		Trr	Pro	Leu	ı Glv	Thr	Ala	. Ala	Cve	: Arc	T Dhe	The same	· '''123	r Phe	T 011	Т~~	
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		Val	Ser	· ጥህተ	Ser	Ser	์ ตา	Len	Pho	TO		. 27-		Leu			
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		7-9	130	neu	Leu	ATA	neu	135	Pro	HIS	Trp	Tyr	Pro 140	Gly	His	Arg	Pro
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		145	Arg	Leu	PIO	Leu	1rp		Cys	Ala	Gly	Val 155		Val	Leu	Ala	Thr
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15		Leu	Pne	Ser	Val	Pro 165	Trp	Leu	Val	Phe	Pro 170		Ala	Ala	Val	Trp	
						•		• • •				*					
		Tyr	Asp	Leu	Val 180	Ile	Cys	Leu	Asp	Phe 185		Asp	Ser	Glu		Leu	Ser
	÷		•	2. 1.				•						•	190	¥	* .
		Leu	Arg	Met 195	Leu	Glu	Val	Leu	Gly 200	Gly	Phe	Leu	Pro	Phe	Leu	Leu	Leu
	,										• . •			205			
20		Leu	Val 210	Cys	His	Val	Leu	Thr	Gln	Ala	Thr	Arg		Суз	His	Arg	Gln
		1,0	210					215			٠.		220	٠		• .	,
		Gln	Gln	Pro	Ala	Ala	Cys	Arg	Gly	Phe	Ala		Val	Ala	Arg.	Thr	Ile
		225		:			230				-	235		*	. *	•	240
25		Leu	Ser	Ala	Tyr	Val	Val	Leu	Arg	Leu	Pro	Tyr	Gln	Leu	Ala	Gln	Leu
23						245					250				٠.	255	-
	* .	Leu	Tyr	Leu	Ala	Phe	Leu	Trp	Asp	Val	Tyr	Ser	Gly	Tyr	Leu	Leu	Trp
*					260				٠	265					270		
		Glu	Ala	Leu	Val	Tyr	Ser	Asp	Tyr	Leu	lle	Leu	Leu	Asn	Ser	Сув	Leu
				275					280			• .		285		įŝ	·-
30		Ser	Pro	Phe	Leu	Cys	Leu	Met	Ala	Ser	Ala	 Asp	Leu	Arg	Thr	Leu	Leu
		• .	290	ŧ				295		٠.			300		. +		
		Arg	Ser	Val	Leu	Ser	Ser	Phe	Ala	Ala	Ala	Leu	Cys	Glu	Glu	Arg	Pro
• 2		305					310	•	-		-	315			<i>:</i>		320
		Gly	Ser	Phe	Thr	Pro	Thr	Glu	Pro	Gln	Thr	Gln	Leu	Asp	Ser	Glu	G] v
35		يد جا		,		325				٠, .	330					335	<i>1</i>
•	•	Pro	Thr	Leu	Pro	Glu	Pro	Met	Ala	Glu	Ala	Gln	Ser	Gln	Met	Asn	Pro
					340					345					250	p	

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Val Ala Gln Pro Gln Val Asn Pro Thr Leu Gln Pro Arg Ser Asp Pro 355 360 365

Thr Ala Gln Pro Gln Leu Asn Pro Thr Ala Gln Pro Gln Ser Asp Pro

5 Thr Ala Gln Pro Gln Leu Asn Leu Met Ala Gln Pro Gln Ser Asp Ser 385

Val Ala Gln Pro Gln Ala Asp Thr Asn Val Gln Thr Pro Ala Pro Ala 405 410 415

Ala Ser Ser Val Pro Ser Pro Cys Asp Glu Ala Ser Pro Thr Pro Ser
420
425
430

Ser His Pro Thr Pro Gly Ala Leu Glu Asp Pro Ala Thr Pro Pro Ala 435 440 445

Ser Glu Gly Glu Ser Pro Ser Ser Thr Pro Pro Glu Ala Ala Pro Gly
450
455
460

15 Ala Gly Pro Thr

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#### (12) INFORMATION FOR SEQ ID NO:11:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1248 base pairs
- 20 (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
  - (ii) MOLECULE TYPE: DNA (genomic)
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

25 ATGTCAGGGA TGGAAAAACT TCAGAATGCT TCCTGGATCT ACCAGCAGAA ACTAGAAGAT 60
CCATTCCAGA AACACCTGAA CAGCACCGAG GAGTATCTGG CCTTCCTCTG CGGACCTCGG 120
CGCAGCCACT TCTTCCTCCC CGTGTCTGTG GTGTATGTGC CAATTTTTGT GGTGGGGGTC 180
ATTGGCAATG TCCTGGTGTG CCTGGTGATT CTGCAGCACC AGGCTATGAA GACGCCCACC 240
AACTACTACC TCTTCAGCCT GGCGGTCTCT GACCTCCTGG TCCTGCTCCT TGGAATGCCC 300
30 CTGGAGGTCT ATGAGATGTG GCGCAACTAC CCTTTCTTGT TCGGGCCCGT GGGCTGCTAC 360
TTCAAGACGG CCCTCTTTGA GACCGTGTGC TTCGCCTCCA TCCTCAGCAT CACCACCGTC 420
AGCGTGGAGC GCTACGTGGC CATCCTACAC CCGTTCCGCG CCAAACTGCA GAGCACCCGG 480
CGCCGGGCCC TCAGGATCCT CGGCATCGTC TGGGGCTTCT CCCCTGCCC 540

AACACCAGCA TCCATGGCAT CAAGTTCCAC TACTTCCCCA ATGGGTCCCT GGTCCCAGGT 600

TCGGCCACCT GTACGGTCAT CAAGCCCATG TGGATCTACA ATTTCATCAT CCAGGTCACC 660

TCCTTCCTAT TCTACCTCCT CCCCATGACT GTCATCAGTG TCCTCTACTA CCTCATGGCA 720

CTCAGACTAA AGAAAGACAA ATCTCTTGAG GCAGATGAAG GGAATGCAAA TATTCAAAGA 780

5 CCCTGCAGAA AATCAGTCAA CAAGATGCTG TTTGTCTTGG TCTTAGTGTT TGCTATCTGT 840

TGGGCCCCGT TCCACATTGA CCGACTCTTC TTCAGCTTTG TGGAGGAGTG GAGTGAATCC 900

CTGGCTGCTG TGTTCAACCT CGTCCATGTG GTGTCAGGTG TCTTCTTCTA CCTGAGCTCA 960

GCTGTCAACC CCATTATCTA TAACCTACTG TCTCGCCGCT TCCAGGCAGC ATTCCAGAAT1020

GTGATCTCTT CTTTCCACAA ACAGTGGCAC TCCCAGCATG ACCCACAGTT GCCACCTGCC1080

10 CAGCGGAACA TCTTCCTGAC AGAATGCCAC TTTGTGGAGC TGACCGAAGA TATAGGTCCC1140

CAATTCCCAT GTCAGTCATC CATGCACAAC TCTCACCTCC CAACAGCCCT CTCTAGTGAA1200

CAGATGTCAA GAACAAACTA TCAAAGCTTC CACTTTAACA AAACCTGA 1248

# (13) INFORMATION FOR SEQ ID NO:12:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 415 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS:
  - (D) TOPOLOGY: not relevant
- (ii) MOLECULE TYPE: protein
- 20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

Met Ser Gly Met Glu Lys Leu Gln Asn Ala Ser Trp Ile Tyr Gln Gln

10
15

Lys Leu Glu Asp Pro Phe Gln Lys His Leu Asn Ser Thr Glu Glu Tyr
20 25 30

Leu Ala Phe Leu Cys Gly Pro Arg Arg Ser His Phe Phe Leu Pro Val

Ser Val Val Tyr Val Pro Ile Phe Val Val Gly Val Ile Gly Asn Val 50 60

Leu Val Cys Leu Val Ile Leu Gln His Gln Ala Met Lys Thr Pro Thr 30 65 70 75 80

Asn Tyr Tyr Leu Phe Ser Leu Ala Val Ser Asp Leu Leu Val Leu Leu 85 90 95

	ьег	1 G17	Me	100	o ren	ı GII	ı Va.	1 Туг	Gl:		: Tr	o Arg	J Ası	1 Ty:	r Pro	) Phe
	Lev	ı Phe	Gly	y Pro	o Val	Gly	γ Cy:	120		e Lys	Thi	Ala	Let 125		e Glu	ı Thr
5	Val	. Cys	Phe	a Ala	l Ser	Ile	Let 135	ı Ser	Ile	Thr	Thi	Va]		· Va]	. Gli	ı Arg
	Tyr 145	Val	Ala	Ile	Leu	His 150	Pro	Phe	: Arg	Ala	Lys 155		ı Gln	Ser	Thr	Arg
10	Arg	Arg	Ala	ı Lev	Arg	Ile	Lev	Gly	Ile	Val 170		Gly	Phe	Ser	Val 175	
	Phe	Ser	Leu	Pro 180	Asn	Thr	Ser	Ile	His 185		Ile	Lys	Phe	His		Phe
	Pro	Asn	Gly 195	Ser	Leu	Val	Pro	Gly 200	Ser	Ala	Thr	Cys	Thr 205	Val	Ile	Lys
15	Pro	Met 210	Trp	Ile	Туг	Asn	Phe 215	Ile	Ile	Gln	Val	Thr 220	Ser	Phe	Leu	Phe
	Tyr 225	Leu	Leu	Pro	Met	Thr 230	Val	Ile	Ser	Val	Leu 235	Tyr	Tyr	Leu	Met	Ala 240
20	Leu	Arg	Leu	Lys	Lys 245	Asp	Lys	Ser	Leu	Glu 250	Ala	Asp	Glu	Gly	Asn 255	
	Asn	Ile	Gln	Arg 260	Pro	Cys	Arg	Lys	Ser 265	Val	Asn	Lys	Met	Leu 270	. ,	Val
	Leu	Val	Leu 275	Val	Phe	Ala	Ile	Cys		Ala	Pro	Phe	_		Asp	Arg
25	Leu	Phe 290		Ser	Phe	Val	Glu 295	Glu	Trp	Ser	Glu		285 Leu	Ala	Ala	Val
	Phe	Asn	Leu	Val	His	Val 310		Ser	Gly	Val		300 Phe	Тут	Leu	Ser	
30			Asn	Pro	Ile		Tyr	Asn	Leu					Phe	Gln	320 Ala
	Ala	Phe	Gln	Asn 340	325 Val	Ile	Ser	Ser	Phe	His	Lys			His		Gln
3.	His	Asp	Pro 355		Leu	Pro	Pro	Ala 360	345 Gln	Arg	Asn	Ile	Phe 365	350 Leu	Thr	Glu
35	Суз	His 370	Phe	Val	Glu	Leu	Thr 375	Glu		Ile		Pro 380		Phe	Pro	Cys
	Gln 385	Ser	Ser	Met	His	Asn 390	Ser	His	Leu		Thr 395	Ala	Leu	Ser	Ser	Glu 400

Gln Met Ser Arg Thr Asn Tyr Gln Ser Phe His Phe Asn Lys Thr 405 410 415

## (14) INFORMATION FOR SEQ ID NO:13:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1173 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- 10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

ATGCCAGATA CTAATAGCAC AATCAATTTA TCACTAAGCA CTCGTGTTAC TTTAGCATTT 60 TTTATGTCCT TAGTAGCTTT TGCTATAATG CTAGGAAATG CTTTGGTCAT TTTAGCTTTT 120 GTGGTGGACA AAAACCTTAG ACATCGAAGT AGTTATTTTT TTCTTAACTT GGCCATCTCT 180 GACTTCTTTG TGGGTGTGAT CTCCATTCCT TTGTACATCC CTCACACGCT GTTCGAATGG 240 15 GATTTTGGAA AGGAAATCTG TGTATTTTGG CTCACTACTG ACTATCTGTT ATGTACAGCA 300 TCTGTATATA ACATTGTCCT CATCAGCTAT GATCGATACC TGTCAGTCTC AAATGCTGTG 360 TCTTATAGAA CTCAACATAC TGGGGTCTTG AAGATTGTTA CTCTGATGGT GGCCGTTTGG 420 GTGCTGGCCT TCTTAGTGAA TGGGCCAATG ATTCTAGTTT CAGAGTCTTG GAAGGATGAA 480 GGTAGTGAAT GTGAACCTGG ATTTTTTCG GAATGGTACA TCCTTGCCAT CACATCATTC 540 20 TTGGAATTCG TGATCCCAGT CATCTTAGTC GCTTATTTCA ACATGAATAT TTATTGGAGC 600 CTGTGGAAGC GTGATCATCT CAGTAGGTGC CAAAGCCATC CTGGACTGAC TGCTGTCTCT 660 TCCAACATCT GTGGACACTC ATTCAGAGGT AGACTATCTT CAAGGAGATC TCTTTCTGCA 720 TCGACAGAAG TTCCTGCATC CTTTCATTCA GAGAGACAGA GGAGAAAGAG TAGTCTCATG 780 TTTTCCTCAA GAACCAAGAT GAATAGCAAT ACAATTGCTT CCAAAATGGG TTCCTTCTCC 840 25 CAATCAGATT CTGTAGCTCT TCACCAAAGG GAACATGTTG AACTGCTTAG AGCCAGGAGA 900 TTAGCCAAGT CACTGGCCAT TCTCTTAGGG GTTTTTGCTG TTTGCTGGGC TCCATATTCT 960 CTGTTCACAA TTGTCCTTTC ATTTTATTCC TCAGCAACAG GTCCTAAATC AGTTTGGTAT1020 AGAATTGCAT TTTGGCTTCA GTGGTTCAAT TCCTTTGTCA ATCCTCTTTT GTATCCATTG1080 TGTCACAAGC GCTTTCAAAA GGCTTTCTTG AAAATATTTT GTATAAAAAA GCAACCTCTA1140 30 CCATCACAAC ACAGTCGGTC AGTATCTTCT TAA 1173

## (15) INFORMATION FOR SEQ ID NO:14:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 390 amino acids
  - (B) TYPE: amino acid
- 5 --- (C) STRANDEDNESS:
  - (D) TOPOLOGY: not relevant
  - (ii) MOLECULE TYPE: protein
    - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:
- Met Pro Asp Thr Asn Ser Thr Ile Asn Leu Ser Leu Ser Thr Arg Val 10 1 5 10
  - Thr Leu Ala Phe Phe Met Ser Leu Val Ala Phe Ala Ile Met Leu Gly
    20 25 30
  - Asn Ala Leu Val Ile Leu Ala Phe Val Val Asp Lys Asn Leu Arg His
    35 40 45
- 15 Arg Ser Ser Tyr Phe Phe Leu Asn Leu Ala Ile Ser Asp Phe Phe Val 50 55 60
  - Gly Val Ile Ser Ile Pro Leu Tyr Ile Pro His Thr Leu Phe Glu Trp
    65 70 75 80
- Asp Phe Gly Lys Glu Ile Cys Val Phe Trp Leu Thr Thr Asp Tyr Leu 20 85 90 95
  - Leu Cys Thr Ala Ser Val Tyr Asn Ile Val Leu Ile Ser Tyr Asp Arg
    100 105 110
  - Tyr Leu Ser Val Ser Asn Ala Val Ser Tyr Arg Thr Gln His Thr Gly
    115 120 125
- Val Leu Lys Ile Val Thr Leu Met Val Ala Val Trp Val Leu Ala Phe
  130
  135
  140
  - Leu Val Asn Gly Pro Met Ile Leu Val Ser Glu Ser Trp Lys Asp Glu
    145 150 155 160
- Gly Ser Glu Cys Glu Pro Gly Phe Phe Ser Glu Trp Tyr Ile Leu Ala 30 165 170 175
  - Ile Thr Ser Phe Leu Glu Phe Val Ile Pro Val Ile Leu Val Ala Tyr
    180 185
  - Phe Asn Met Asn Ile Tyr Trp Ser Leu Trp Lys Arg Asp His Leu Ser 195 200 205
- Arg Cys Gln Ser His Pro Gly Leu Thr Ala Val Ser Ser Asn Ile Cys 210 215 220

	22	у ніз 5	s Ser	Pne	: Arg	230	/ Arg	, Lei	ı Ser	Ser	Arg 235		Ser	Leu	Ser	Ala 240
	Sea	r Thi	Glu	Val	Pro 245	Ala	Ser	Phe	His	Ser 250	Glu	Arg	Gln	Arg	Arg 255	
5	Ser	Ser	Leu	Met 260	Phe	Ser	Ser	Arg	Thr 265	Lys	Met	Asn	Ser	Asn 270		Ile
	Ala	Ser	Lys 275	Met	Gly	Ser	Phe	Ser 280	Gln	Ser	Asp		Val 285	Ala	Leu	His
10	Gln	Arg 290	Glu	His	Val	Glu	Leu 295	Leu	Arg	Ala	Arg	Arg 300	Leu	Ala	Lys	Ser
	Leu 305	Ala	Ile	Leu	Leu	Gly 310	Val	Phe	Ala	Val	Cys 315		Ala	Pro	Tyr	Ser 320
	Leu	Phe	Thr	Ile	Val 325	Leu	Ser	Phe	Tyr	Ser 330	Ser	Ala	Thr	Gly	Pro 335	Lys
15	Ser	Val	Trp	Tyr 340	Arg	Ile	Ala	Phe	Trp 345	Leu	Gln	Trp		Asn 350	Ser	Phe
*	Val	Asn	Pro 355	Leu	Leu	Tyr	Pro	Leu 360	Cys	His	Lys	Arg	Phe	Gln	Lys	Ala
20	Phe	Leu 370	Lys	Ile	Phe	Cys	Ile 375	Lys	Lys	Gln		Leu 380		Ser	Gln	His
	Ser 385	Arg	Ser	Val		Ser 390						* 1				
(16)			CION						*	45					2 - 14 2	
25	(i)	(A) (B) (C)	LEN TYP STR TOP	GTH: E: n ANDE	1120 ucle: DNES	B ba ic a B: s	se p cid ingl	airs				1				
*	(ii)	MOLE	CULE	TYP	E: DI	VA (	geno	mic)								

# 30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:

ATGGCGAACG CGAGCGAGCC GGGTGGCAGC GGCGGCGGC AGGCGGCCGC CCTGGGCCTC 60

AAGCTGGCCA CGCTCAGCCT GCTGCTGTC GTGAGCCTAG CGGGCAACGT GCTGTTCGCG 120

CTGCTGATCG TGCGGGAGCG CAGCCTGCAC CGCGCCCCGT ACTACCTGCT GCTCGACCTG 180

TGCCTGGCCG ACGGCTGCG CGCGCCCCG TGCCTCCCGG CCGTCATGCT GGCGGCGCG 240

35 CGTGCGGCGG CCGCGGG GGCGCCGC GGCGCCCGG GCTGCAAGCT GCTCGCCTTC 300

CTGGCCGCC TCTTCTGCTT CCACGCCGCC TTCCTGCTGC TGGGCGTGGG CGTCACCCGC 360

TACCTGGCCA TCGCGCACCA CCGCTTCTAT GCAGAGCGCC TGGCCGGCTG GCCGTGCGCC 420

GCCATGCTGG TGTGCGCCGC CTGGGCGCTG GCGCTGGCCG CGGCCTTCCC GCCAGTGCTG 480

GACGGCGGTG GCGACGACGA GGACGCGCG TGCGCCCTGG AGCAGCGGCC CGACGGCGCC 540

5 CCCGGCGGCG TGGGCTTCCT GCTGCTGCTG GCCGTGGTGG TGGGCCCACC GCACCTCGTC 600

TACCTCCGCC TGCTCTTCTT CATCCACGAC CGCCGCAAGA TGCGGCCCAC GCGCCTGGTG 660

CCCGCCGTCA GCCACGACTG GACCTTCCAC GGCCCGGGCG CCACCGGCCA GGCGGCCGC 720

AACTGGACGG CGGGCTTCGG CCGCGGGCCC ACGCCGCCG CGCTTGTGGG CATCCGGCCC 780

GCAGGGCCGG GCCGCGGCG GCGCCGCCC CTCGTGCTGG AAGAATTCAA GACGGAGAAG 840

10 AGGCTGTGCA AGATGTTCTA CGCCGTCACG CTGCTCTTCC TGCTCCTCTG GGGGCCCTAC 900

GTCGTGGCCA GCTACCTGCG GGTCCTGGTG CGGCCCGGCG CCGTCCCCCA GGCCTACCTG 960

ACGGCCTCCG TGTGGCTGAC CTTCGCGCAG GCCCGGCATCA ACCCCGTCGT GTGCTTCCTC1020

TTCAACAGGG AGCTGAGGAA CTGCTTCAGG GCCCAGTTCC CCTGCTGCCA GAGCCCCCGG1080

ACCACCCAGG CGACCCATCC CTGCGACCTG AAAGGCATTG GTTTATGA 1128

# 15 (17) INFORMATION FOR SEQ ID NO:16:

20

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 375 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS:
- (D) TOPOLOGY: not relevant
  - (ii) MOLECULE TYPE: protein
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:

Met Ala Asn Ala Ser Glu Pro Gly Gly Ser Gly Gly Gly Glu Ala Ala

1 10 15

Ala Leu Gly Leu Lys Leu Ala Thr Leu Ser Leu Leu Cys Val Ser

20 25 30

Leu Ala Gly Asn Val Leu Phe Ala Leu Leu Ile Val Arg Glu Arg Ser

Leu His Arg Ala Pro Tyr Tyr Leu Leu Asp Leu Cys Leu Ala Asp
50 55 60

Gly Leu Arg Ala Leu Ala Cys Leu Pro Ala Val Met Leu Ala Ala Arg
65 70 75 80

	Arg	, Ala	ı Ala	Ala	Ala 85	Ala	Gly	Ala	Pro	90	Gly	/ Ala	. Leu	Gly	95	Lys
	Leu	ı Leu	Ala	Phe 100		Ala	Ala	Lev	Phe		Phe	His	. Ala	Ala 110		Leu
5	Leu	Leu	Gly 115	Val	Gly	Val	Thr	Arg		Leu	Ala	Ile	Ala 125		His	Arg
	Dho	Mis said	הוא הי	C1.,	7.~~	, tan	 							. 49	****	
*	Pile	130	YIG	Giu	Arg	reu	135		Trp	Pro	Cys	140	Ala	Met	Leu	Val
	Cys	Ala	Ala	Trp	Ala	Leu	Ala	Leu	Ala	Ala	Ala	Phe	Pro	Pro	Val	Leu
10	145					150			,	• • •	155					160
:	Asp	Gly	Gly	Gly	Asp	Asp	Glu	Asp	Ala		Cys	Ala	Leu	Glu		Arg
				13.				* :	*	170	:	• •		Ý	175	
	Pro	Asp	Gly	Ala 180		Gly	Ala	Leu	Gly 185		Leu	Leu	Leu	Leu 190	Ala	Val
15	Val	Val	Gly 195	Ala	Thr	His	Leu		Tyr	Leu	Arg	Leu	Leu	Phe	Phe	Ile
		*		+ 1	*		-	. 200					205	:		
	His	Asp 210	Arg	Arg	Lys	Met	Arg 215	Pro	Ala	Arg	Leu	Val 220	Pro	Ala	Val	Ser
20	His 225		Trp	Thr	Phe	His 230	Gly	Pro	Gly	Ala	Thr 235	Gly	Gln	Ala	Ala	
			mi				·.			_			,	- 10		240
(9)	ASII	irp	Inr	Ala	245	Pne	GIA	Arg	GIA	250	Thr	Pro	Pro	Ala	Leu 255	Val.
	Gly	Ile	Arg	Pro 260	Ala	Gly	Pro	Gly	Arg 265	Gly	Ala	Arg	Arg	Leu 270		Val
25	Leu	Glu	Glu 275	Phe	Lys	Thr	Glu	Lys 280	Arg	Leu	Сув	Lys	Met 285	Phe	Tyr	Ala
	Val	Thr 290	Leu	Leu	Phe	Leu	Leu 295	Leu	Trp	Gly	Pro	Tyr 300	Val	Val	Ala	Ser
30	Tyr 305	Leu	Arg	Val	Leu	Val 310	Arg	Pro	Gly	Ala	Val 315	Pro	Gln	Ala	Tyr	Leu 320
	Thr	Ala	Ser	Val	Trp 325	Leu	Thr	Phe	Ala	Gln 330	Ala	Gly	Ile	Asn	Pro 335	Val
•	Val	Cys		Leu 340	Phe	Asn	Arg	Glu	Leu 345	Arg	Asp	Cys	Phe	Arg 350		Gln
35	Phe	Pro	Cys 355	Cys	Gln	Ser	Pro	Arg 360	Thr	Thr	Gln	Ala	Thr 365	His	Pro	Сув
r Conten	n en	Tou	T 110	C11-	T1 a'	<b>.</b>	<b>V</b>		: · · ·			- -: .				

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# (18) INFORMATION FOR SEQ ID NO:17:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1002 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:
- 10 ATGAACACCA CAGTGATGCA AGGCTTCAAC AGATCTGAGC GGTGCCCCAG AGACACTCGG 60 ATAGTACAGC TGGTATTCCC AGCCCTCTAC ACAGTGGTTT TCTTGACCGG CATCCTGCTG 120 AATACTTTGG CTCTGTGGGT GTTTGTTCAC ATCCCCAGCT CCTCCACCTT CATCATCTAC 180 CTCAAAAACA CTTTGGTGGC CGACTTGATA ATGACACTCA TGCTTCCTTT CAAAATCCTC 240 TCTGACTCAC ACCTGGCACC CTGGCAGCTC AGAGCTTTTG TGTGTCGTTT TTCTTCGGTG 300 15 ATATTTTATG AGACCATGTA TGTGGGCATC GTGCTGTTAG GGCTCATAGC CTTTGACAGA 360 TTCCTCAAGA TCATCAGACC TTTGAGAAAT ATTTTTCTAA AAAAACCTGT TTTTGCAAAA 420 ACGGTCTCAA TCTTCATCTG GTTCTTTTTG TTCTTCATCT CCCTGCCAAA TACGATCTTG 480 AGCAACAAGG AAGCAACACC ATCGTCTGTG AAAAAGTGTG CTTCCTTAAA GGGGCCTCTG 540 GGGCTGAAAT GGCATCAAAT GGTAAATAAC ATATGCCAGT TTATTTTCTG GACTGTTTTT 600 20 ATCCTAATGC TTGTGTTTTA TGTGGTTATT GCAAAAAAAG TATATGATTC TTATAGAAAG 660 TCCAAAAGTA AGGACAGAAA AAACAACAAA AAGCTGGAAG GCAAAGTATT TGTTGTCGTG 720 GCTGTCTTCT TTGTGTGTTT TGCTCCATTT CATTTTGCCA GAGTTCCATA TACTCACAGT 780 CAAACCAACA ATAAGACTGA CTGTAGACTG CAAAATCAAC TGTTTATTGC TAAAGAAACA 840 ACTCTCTTTT TGGCAGCAAC TAACATTTGT ATGGATCCCT TAATATACAT ATTCTTATGT 900 25 AAAAAATTCA CAGAAAAGCT ACCATGTATG CAAGGGAGAA AGACCACAGC ATCAAGCCAA 960 GAAAATCATA GCAGTCAGAC AGACAACATA ACCTTAGGCT GA 1002
  - (19) INFORMATION FOR SEQ ID NO:18:
    - (i) SEQUENCE CHARACTERISTICS:
      - (A) LENGTH: 333 amino acids
    - (B) TYPE: amino acid
      - (C) STRANDEDNESS:

35

- (D) TOPOLOGY: not relevant
- (ii) MOLECULE TYPE: protein

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	•	· (XI)	SEC	OENC	E DE	SCRI	PTIC	ON: S	EQ 1	LD NC	):18:					٠, .	
5		Met	Asn	Thr	Thr	Val	Met	Gln	Gly	Phe	Asr	Arg	Ser	Glu	Arg	Суз	Pro
5	1	1			-	5					10					15	
	•	Arg	Asp	Thr	Arg 20	Ile	Val	Gln	Leu	Val 25	Phe	Pro	Ala	Leu	Tyr 30	Thr	Val
		Val	Phe	Leu 35	Thr	Gly	Ile	Leu	Leu 40	Asn	Thr	Leu	Ala	Leu 45	Trp	Val	Phe
10		Val	His 50	Ile	Pro	Ser	Ser	Ser 53	Thr	Phe	Ile	Ile	Tyr 60	Leu	Lys	Asn	Thr
		Leu 65	Val	Ala	Asp	Leu	Ile	Met	Thr	Leu	Met		Pro	Phe	Lys	Ile	Leu
				1						٠.	•	75		٠			80
15		Ser	Asp	Ser	His	Leu 85	Ala	Pro	Trp	Gln	Leu 90	Arg	Ala	Phe	Val	Cys 95	Arg
		Phe	Ser	Ser	Val	Ile	Phe	Tyr	Glu	Thr 105	Met	Туг	Val	Gly	Ile 110	Val	Leu
		Leu	Gly	Leu 115	Ile	Ala	Phe	Asp	Arg 120	Phe	Leu	Lys	Ile	Ile 125	Arg	Pro	Leu
20	•	Arq	Asn	Ile	Phe	Leu	Lvs	Lvs	Pro	Val	Dhe	בות	Tara		17.1		Ile.
			130				, T	135			1116	AIG.	140	IIII	val	Ser	TTG.
		Phe	Ile	Trp	Phe	Phe	Leu	Phe	Phe	Ile	Ser	Leu	Pro	Asn	Thr	Ile	Leu
		143					150				•	155	-	* .		:	160
25		Ser	Asn	Lys	Glu	Ala 165	Thr	Pro	Ser	Ser	Val 170	Lys	Lys	Cys		Ser 175	Leu
	•	Lys	Gly	Pro	Leu 180	Gly	Leu		Trp	300		Met	Val		Asn 190		Cys
		Gln	Phe	Ile 195	Phe	Trp	Thr	Val	Phe	Ile	Leu	Met	Leu	Val	Phe		Val
20									200	•				205			
30 .		Val	Ile 210	Ala	Lys	Lys	Val	Tyr 215	Asp	Ser	Tyr	Arg	Lys 220	Ser	Lys	Ser	Lys
•		Asp 225	Arg	Lys	Asn	Asn	Lys 230	Lys	Leu	Glu	Gly	Lys	Val	Phe	Val	Val	Val

230

245

Ala Val Phe Phe Val Cys Phe Ala Pro Phe His Phe Ala Arg Val Pro

- Tyr Thr His Ser Gln Thr Asn Asn Lys Thr Asp Cys Arg Leu Gln Asn 260 265 270
- Gln Leu Phe Ile Ala Lys Glu Thr Thr Leu Phe Leu Ala Ala Thr Asn
  275
  280
  285
- Ile Cys Met Asp Pro Leu Ile Tyr Ile Phe Leu Cys Lys Lys Phe Thr 290 295 300

Glu Lys Leu Pro Cys Met Gln Gly Arg Lys Thr Thr Ala Ser Ser Gln 305 310 315

Glu Asn His Ser Ser Gln Thr Asp Asn Ile Thr Leu Gly
325 330

- (20) INFORMATION FOR SEQ ID NO:19:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 1122 base pairs
    - (B) TYPE: nucleic acid
- 15 (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
  - (ii) MOLECULE TYPE: DNA (genomic)
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:
- ATGGCCACA CTACCGGAGA GCCTGAGGAG GTGAGCGGCG CTCTGTCCCC ACCGTCCGCA 60

  20 TCAGCTTATG TGAAGCTGGT ACTGCTGGGA CTGATTATGT GCGTGAGCCT GGCGGGTAAC 120

  GCCATCTTGT CCCTGCTGGT GCTCAAGGAG CGTGCCCTGC ACAAGGCTCC TTACTACTTC 180

  CTGCTGGACC TGTGCCTGGC CGATGGCATA CGCTCTGCCG TCTGCTTCCC CTTTGTGCTG 240

  GCTTCTGTGC GCCACGGCTC TTCATGGACC TCAGGTGCAA GATTGTGGCC 300

  TTTATGGCCG TGCTCTTTG CTTCCATGCG GCCTTCATGC TGTTCTGCAT CAGCGTCACC 360

  25 CGCTACATGG CCATCGCCCA CCACCGCTTC TACGGCAAGC GCATGACACT CTGGACATGC 420

  GCGGCTGTCA TCTGCATGGC CTGGACCCTG TCTGTGGCCA TGGCCTTCCC ACCTGTCTT 480

  GACGTGGGCA CCTACAAGTT TATTCGGGAG GAGGACCAGT GCATCTTTGA GCATCGCTAC 540

  TTCAAGGCCA ATGACACGCT GGGCTTCATG CTTATGTTGG CTGTGCTCAT GGCAGCTACC 600

  CATGCTGTCT ACGGCAAGCT GCTCCTCTC GAGTATCGTC ACCGCAAGAT GAAGCCAGTG 660

  30 CAGATGGTGC CAGCCATCAG CCAGAACTGG ACATTCCATG GTCCCGGGGC CACCGGCCAG 720

  GCTGCTGCCA ACTGGATCGC CGGCTTTGGC CGTGGGCCCA TGCCCAACC CCTGCTGGGT 780

ATCCGGCAGA ATGGGCATGC AGCCAGCCGG CGGCTACTGG GCATGGACGA GGTCAAGGGT 840

GAAAAGCAGC TGGGCCGCAT GTTCTACGCG ATCACACTGC TCTTTCTGCT CCTCTGGTCA 900

CCCTACATCG TGGCCTGCTA CTGGCGAGTG TTTGTGAAAG CCTGTGCTGT GCCCCACCGC 960

TACCTGGCCA CTGCTGTTTG GATGAGCTTC GCCCAGGCTG CCGTCAACCC AATTGTCTGC1020

TTCCTGCTCA ACAAGGACCT CAAGAAGTGC CTGACCACTC ACGCCCCCTG CTGGGGCACA1080

5 GGAGGTGCCC CGGCTCCCAG AGAACCCTAC TGTGTCATGT GA 1122

## (21) INFORMATION FOR SEQ ID NO:20:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 373 amino acids
  - (B) TYPE: amino acid
- 10
- (C) STRANDEDNESS:
- (D) TOPOLOGY: not relevant
- (ii) MOLECULE TYPE: DNA (genomic)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:
- Met Ala Asn Thr Thr Gly Glu Pro Glu Glu Val Ser Gly Ala Leu Ser

  15 1 5 10 15
  - Pro Pro Ser Ala Ser Ala Tyr Val Lys Leu Val Leu Leu Gly Leu Ile 20 25 30
  - Met Cys Val Ser Leu Ala Gly Asn Ala Ile Leu Ser Leu Leu Val Leu 35 40 45
- Lys Glu Arg Ala Leu His Lys Ala Pro Tyr Tyr Phe Leu Leu Asp Leu
  50 55 60
  - Cys Leu Ala Asp Gly Ile Arg Ser Ala Val Cys Phe Pro Phe Val Leu 65 70 75 80
- Ala Ser Val Arg His Gly Ser Ser Trp Thr Phe Ser Ala Leu Ser Cys
  85 90 95
  - Lys Ile Val Ala Phe Met Ala Val Leu Phe Cys Phe His Ala Ala Phe
    100 105 110
  - Met Leu Phe Cys Ile Ser Val Thr Arg Tyr Met Ala Ile Ala His His
    115 120 125
- Arg Phe Tyr Ala Lys Arg Met Thr Leu Trp Thr Cys Ala Ala Val Ile
  130 135 140
  - Cys Met Ala Trp Thr Leu Ser Val Ala Met Ala Phe Pro Pro Val Phe 145 150 155 160
  - Asp Val Gly Thr Tyr Lys Phe Ile Arg Glu Glu Asp Gln Cys Ile Phe 165 170 175

	Glı	u Hi	s Ar	g Tyı 180	r Pho	e Ly	s Ala	a Ası	n Asp 185	Thi	Lei	ı Gl	y Phe	Mei 19		u Met
	Let	u Al	a Va 19	l Leu 5	ı Met	t Ala	a Ala	200	r His	3 Ala	va]	. Туз	Cly 205		s Let	ı Lev
5	Let	1 Pho 210	e Glu	1 Туг	Arg	g His	215	J Lys	3 Met	. Lys	Pro	Va]	L Gln	Met	. Val	Pro
	Ala 225	ı Ile	e Ser	Gln	Asr	230	Thr	Phe	: His	Gly	Pro 235	Gly	/ Ala	Thr	Gly	Gln 240
10	Ala	Ala	a Ala	Asn	Trp 245	Ile	Ala	Gly	Phe	Gly 250	Arg	Gly	Pro	Met	Pro 255	
	Thr	Leu	Leu	Gly 260	Ile	Arg	Gln	Asn	Gly 265	His	Ala	Ala	Ser	Arg 270	Arg	Leu
	Leu	Gly	Met 275	Asp	Glu	Val	Lys	Gly 280	Glu	Lys	Gln	Leu	Gly 285	Arg	Met	Phe
15	Tyr	Ala 290	Ile	Thr	Leu	Leu	Phe 295	Leu	Leu	Leu	Trp	Ser 300	Pro	Tyr	Ile	Val
	Ala 305	Cys	Tyr	Trp	Arg	Val 310	Phe	Val	Lys	Ala	Cys 315	Ala	Val	Pro	His	Arg 320
20	Tyr	Leu	Ala	Thr	Ala 325	Val	Trp	Met	Ser	Phe 330	Ala	Gln	Ala	Ala	Val 335	Asn
	Pro	Ile	Val	Cys 340	Phe	Leu	Leu	Asn	Lys 345	Asp	Leu	Lys	Lys	Суs 350	Leu	Thr
	Thr	His	Ala 355	Pro	Суз	Trp	Gly	Thr 360	Gly	Gly	Ala	Pro	Ala 365	Pro	Arg	Glu
25	Pro	Tyr 370	Cys	Val	Met	*		e For		* * * * * * * * * * * * * * * * * * * *			1 - ja	*		
(22)	INFO	RMAT	NOI	FOR	SEQ	ID N	0:21	• •	· · i.		*		· .		n	
• • • •	(i)	SEQU	JENCE	CHA	RACT	ERIS	TICS	:				` . ; ·	. *			. :

- (A) LENGTH: 1053 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:
- 35 ATGGCTTTGG AACAGAACCA GTCAACAGAT TATTATTATG AGGAAAATGA AATGAATGGC 60
  ACTTATGACT ACAGTCAATA TGAATTGATC TGTATCAAAG AAGATGTCAG AGAATTTGCA 120

AAAGTTTTCC TCCCTGTATT CCTCACAATA GCTTTCGTCA TTGGACTTGC AGGCAATTCC 180 ATGGTAGTGG CAATTTATGC CTATTACAAG AAACAGAGAA CCAAAACAGA TGTGTACATC 240 CTGAATTTGG CTGTAGCAGA TTTACTCCTT CTATTCACTC TGCCTTTTTG GGCTGTTAAT 300 GCAGTTCATG GGTGGGTTTT AGGGAAAATA ATGTGCAAAA TAACTTCAGC CTTGTACACA 360 5 CTAAACTTTG TCTCTGGAAT GCAGTTTCTG GCTTGCATCA GCATAGACAG ATATGTGGCA 420 GTAACTAATG TCCCCAGCCA ATCAGGAGTG GGAAAACCAT GCTGGATCAT CTGTTTCTGT 480 GTCTGGATGG CTGCCATCTT GCTGAGCATA CCCCAGCTGG TTTTTTATAC AGTAAATGAC 540 AATGCTAGGT GCATTCCCAT TTTCCCCCGC TACCTAGGAA CATCAATGAA AGCATTGATT 600 CAAATGCTAG AGATCTGCAT TGGATTTGTA GTACCCTTTC TTATTATGGG GGTGTGCTAC 660 10 TTTATCACGG CAAGGACACT CATGAAGATG CCAAACATTA AAATATCTCG ACCCTAAAA 720 GTTCTGCTCA CAGTCGTTAT AGTTTTCATT GTCACTCAAC TGCCTTATAA CATTGTCAAG 780 TTCTGCCGAG CCATAGACAT CATCTACTCC CTGATCACCA GCTGCAACAT GAGCAAACGC 840 ATGGACATCG CCATCCAAGT CACAGAAAGC ATTGCACTCT TTCACAGCTG CCTCAACCCA 900 ATCCTTTATG TTTTTATGGG AGCATCTTTC AAAAACTACG TTATGAAAGT GGCCAAGAAA 960 15 TATGGGTCCT GGAGAAGACA GAGACAAAGT GTGGAGGAGT TTCCTTTTGA TTCTGAGGGT1020 CCTACAGAGC CAACCAGTAC TTTTAGCATT TAA 1053

- (23) INFORMATION FOR SEQ ID NO:22:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 350 amino acids
- 20 (B) TYPE: amino acid
  - (C) STRANDEDNESS:
  - (D) TOPOLOGY: not relevant
  - (ii) MOLECULE TYPE: protein
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:
- Met Ala Leu Glu Gln Asn Gln Ser Thr Asp Tyr Tyr Tyr Glu Glu Asn

  1 5 10 15
  - Glu Met Asn Gly Thr Tyr Asp Tyr Ser Gln Tyr Glu Leu Ile Cys Ile
    20 25 30
- Lys Glu Asp Val Arg Glu Phe Ala Lys Val Phe Leu Pro Val Phe Leu 35 40 45
  - Thr Ile Ala Phe Val Ile Gly Leu Ala Gly Asn Ser Met Val Val Ala

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	. Il 65	е Ту	r Al	а Ту	т Ту	70	s Ly	s Gl	n Ar	g Th	r Ly 75	s Th	r Asp	va:	l Ty	r I. 80
5	Le	u Ası	n Le	u Al	a Va 85	l Al	a Asj	p Le	u Lei	u -Lei 90	ı Lei	ı Phe	∋ Thr	Lei	1 Pro	o Pl
	Trj	o Ala	a Va	l As 10	n Ala	a Val	L His	s Gl	y Tr 10	o Val	l Lei	ı Gly	/ Lys	Ile 110		Су
	Lys	: Ile	Th:	r Se: 5	r Ala	a Lei	туг	Th:	Let	ı Asr	n Phe	Val	. Ser 125		Met	: G1
10	Phe	Leu 130	l Ala	i Cys	3 Ile	e Ser	Ile 135	Asp	Arg	ј Туг	Val	Ala 140	Val	Thr	Asn	ı. Va
	Pro 145	Ser	Glr	Ser	Gly	/ Val 150	Gly	' Lys	Pro	Cys	Trp	Ile	Ile	Cys	Phe	Су 16
15	Val	Trp	Met	Ala	Ala 165	Ile	Leu	Leu	Ser	Ile 170	Pro	Gln	Leu	Val	Phe	
	Thr	Val	Asn	Asp 180	Asn	Ala	Arg	Суз	Ile 185	Pro	Ile	Phe	Pro	Arg	100	Le
	Gly	Thr	Ser 195	Met	Lys	Ala	Leu	Ile 200	Gln	Met	Leu	Glu	Ile 205		Ile	Gly
20	Phe	Val 210	Val	Pro	Phe	Leu	Ile 215	Met	Gly	Val	Cys	Tyr 220	Phe	Ile	Thr	Ala
	Arg 225	Thr	Leu	Met	Lys	Met 230	Pro	Asn	Ile	Lys	Ile 235		Arg	Pro	Leu	
25	Val	Leu	Leu	Thr	Val 245	Val	Ile	Val	Phe	Ile 250		Thr	Gln	Leu		240 Tyr
	Asn	Ile	Val	Lys 260	Phe	Суз	Arg	Ala	Ile 265	×	Ile	Ile	Tyr		255 Leu	Ile
	Thr	Ser	Cys 275	Asn	Met	Ser	Lys	Arg 280		Asp	Ile		Ile (	270 31n	Val	Thr
30	Glu	Ser 290	Ile	Ala	Leu	Phe			Cys	Leu		Pro	285 Ile 1	Jeu '	Tyr	Val
	Phe 1	Met (	Gly	Ala	Ser			Asn	Tyr	Val 1	Met :	rys ,	Val <i>I</i>	la 1	Lys :	Lys
35	Tyr	Gly :	Ser	Trp			3ln i	Arg	Gln	Ser 1	315 Val (	Glu (	Slu F	he l	· ·	320 Phe
1 42	Asp S			·		. · ' ;				330	* 11			3	335	. i :

#### (24) INFORMATION FOR SEQ ID NO:23:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1116 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:

ATGCCAGGAA ACGCCACCCC AGTGACCACC ACTGCCCCGT GGGCCTCCCT GGGCCTCTCC 60 10 GCCAAGACCT GCAACAACGT GTCCTTCGAA GAGAGCAGGA TAGTCCTGGT CGTGGTGTAC 120 AGCGCGGTGT GCACGCTGGG GGTGCCGGCC AACTGCCTGA CTGCGTGGCT GGCGCTGCTG 180 CAGGTACTGC AGGGCAACGT GCTGGCCGTC TACCTGCTCT GCCTGGCACT CTGCGAACTG 240 CTGTACACAG GCACGCTGCC ACTCTGGGTC ATCTATATCC GCAACCAGCA CCGCTGGACC 300 CTAGGCCTGC TGGCCTCGAA GGTGACCGCC TACATCTTCT TCTGCAACAT CTACGTCAGC 360 15 ATCCTCTTCC TGTGCTGCAT CTCCTGCGAC CGCTTCGTGG CCGTGGTGTA CGCGCTGGAG 420 AGTCGGGGCC GCCGCCGCC GAGGACCGCC ATCCTCATCT CCGCCTGCAT CTTCATCCTC 480 GTCGGGATCG TTCACTACCC GGTGTTCCAG ACGGAAGACA AGGAGACCTG CTTTGACATG 540 CTGCAGATGG ACAGCAGGAT TGCCGGGTAC TACTACGCCA GGTTCACCGT TGGCTTTGCC 600 ATCCCTCTCT CCATCATCGC CTTCACCAAC CACCGGATTT TCAGGAGCAT CAAGCAGAGC 660 20 ATGGGCTTAA GCGCTGCCCA GAAGGCCAAG GTGAAGCACT CGGCCATCGC GGTGGTTGTC 720 ATCTTCCTAG TCTGCTTCGC CCCGTACCAC CTGGTTCTCC TCGTCAAAGC CGCTGCCTTT 780 TCCTACTACA GAGGAGACAG GAACGCCATG TGCGGCTTGG AGGAAAGGCT GTACACAGCC 840 TCTGTGGTGT TTCTGTGCCT GTCCACGGTG AACGGCGTGG CTGACCCCAT TATCTACGTG 900 CTGGCCACGG ACCATTCCCG CCAAGAAGTG TCCAGAATCC ATAAGGGGTG GAAAGAGTGG 960 25 TCCATGAAGA CAGACGTCAC CAGGCTCACC CACAGCAGGG ACACCGAGGA GCTGCAGTCG1020 CCCGTGGCCC TTGCAGACCA CTACACCTTC TCCAGGCCCG TGCACCCACC AGGGTCACCA1080 TGCCCTGCAA AGAGGCTGAT TGAGGAGTCC TGCTGA 1116

# (25) INFORMATION FOR SEQ ID NO:24:

<sup>(</sup>i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 371 amino acids

- (B) TYPE: amino acid
- (C) STRANDEDNESS:
- (D) TOPOLOGY: not relevant
- (ii) MOLECULE TYPE: protein
- 5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:24:

Met Pro Gly Asn Ala Thr Pro Val Thr Thr Thr Ala Pro Trp Ala Ser

1 10 15

Leu Gly Leu Ser Ala Lys Thr Cys Asn Asn Val Ser Phe Glu Glu Ser 20 25 30

10 Arg Ile Val Leu Val Val Val Tyr Ser Ala Val Cys Thr Leu Gly Val
35 40 45

Pro Ala Asn Cys Leu Thr Ala Trp Leu Ala Leu Leu Gln Val Leu Gln 50 55 60

Gly Asn Val Leu Ala Val Tyr Leu Leu Cys Leu Ala Leu Cys Glu Leu 15 65 70 75 80

Leu Tyr Thr Gly Thr Leu Pro Leu Trp Val Ile Tyr Ile Arg Asn Gln 85 90 95

His Arg Trp Thr Leu Gly Leu Leu Ala Ser Lys Val Thr Ala Tyr Ile
100 105 110

20 Phe Phe Cys Asn Ile Tyr Val Ser Ile Leu Phe Leu Cys Cys Ile Ser 115 120 125

Cys Asp Arg Phe Val Ala Val Val Tyr Ala Leu Glu Ser Arg Gly Arg 130 135 140

Arg Arg Arg Thr Ala Ile Leu Ile Ser Ala Cys Ile Phe Ile Leu 25 145 150 155 160

Val Gly Ile Val His Tyr Pro Val Phe Gln Thr Glu Asp Lys Glu Thr
165 170 175

Cys Phe Asp Met Leu Gln Met Asp Ser Arg Ile Ala Gly Tyr Tyr
180 185 190

Ala Arg Phe Thr Val Gly Phe Ala Ile Pro Leu Ser Ile Ile Ala Phe
195 200 205

Thr Asn His Arg Ile Phe Arg Ser Ile Lys Gln Ser Met Gly Leu Ser 210 215 220

Ala Ala Gln Lys Ala Lys Val Lys His Ser Ala Ile Ala Val Val
5 225 230 235 240

Ile Phe Leu Val Cys Phe Ala Pro Tyr His Leu Val Leu Leu Val Lys
245 250 255

可是"是我们的一个"

- Ala Ala Ala Phe Ser Tyr Tyr Arg Gly Asp Arg Asn Ala Met Cys Gly
  260 265 270
- 5 Leu Glu Glu Arg Leu Tyr Thr Ala Ser Val Val Phe Leu Cys Leu Ser 275 280 285
  - Thr Val Asn Gly Val Ala Asp Pro Ile Ile Tyr Val Leu Ala Thr Asp 290 295 300
- His Ser Arg Gln Glu Val Ser Arg Ile His Lys Gly Trp Lys Glu Trp 305 310 315 320
  - Ser Met Lys Thr Asp Val Thr Arg Leu Thr His Ser Arg Asp Thr Glu 325 330 335
  - Glu Leu Gln Ser Pro Val Ala Leu Ala Asp His Tyr Thr Phe Ser Arg 340 345 350
- Pro Val His Pro Pro Gly Ser Pro Cys Pro Ala Lys Arg Leu Ile Glu 355 360 365

Glu Ser Cys

- (26) INFORMATION FOR SEQ ID NO:25:
- 20 (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1113 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- 25 (ii) MOLECULE TYPE: DNA (genomic)
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:25:

ATGGCGAACT ATAGCCATGC AGCTGACAAC ATTTTGCAAA ATCTCTCGCC TCTAACAGCC 60

TTTCTGAAAC TGACTTCCTT GGGTTTCATA ATAGGAGTCA GCGTGGTGGG CAACCTCCTG 120

ATCTCCATTT TGCTAGTGAA AGATAAGACC TTGCATAGAG CACCTTACTA CTTCCTGTTG 180

30 GATCTTTGCT GTTCAGATAT CCTCAGATCT GCAATTTGTT TCCCATTTGT GTTCAACTCT 240

GTCAAAAATG GCTCTACCTG GACTTATGGG ACTCTGACTT GCAAAGTGAT TGCCTTTCTG 300

GGGGTTTTGT CCTGTTTCCA CACTGCTTTC ATGCTCTTCT GCATCAGTGT CACCAGATAC 360

TTAGCTATCG CCCATCACCG CTTCTATACA AAGAGGCTGA CCTTTTGGAC GTGTCTGGCT 420

GTGATCTGTA TGGTGTGGAC TCTGTCTGTG GCCATGGCAT TTCCCCCGGT TTTAGACGTG 480

GGCACTTACT CATTCATTAG GGAGGAAGAT CAATGCACCT TCCAACACCG CTCCTTCAGG 540
GCTAATGATT CCTTAGGATT TATGCTGCTT CTTGCTCTCA TCCTCCTAGC CACACAGCTT 600
GTCTACCTCA AGCTGATATT TTTCGTCCAC GATCGAAGAA AAATGAAGCC AGTCCAGTTT 660
GTAGCAGCAG TCAGCCAGAA CTGGACTTTT CATGGTCCTG GAGCCAGTGG CCAGGCAGCT 720
5 GCCAATTGGC TAGCAGGATT TGGAAGGGGT CCCACACCAC CCACCTTGCT GGGCATCAGG 780
CAAAATGCAA ACACCACAGG CAGAAGAAGG CTATTGGTCT TAGACCAGTT CAAAATGGAG 840
AAAAGAATCA GCAGAATGTT CTATATAATG ACTTTTCTGT TTCTAACCTT GTGGGGCCCC 900
TACCTGGTGG CCTGTTATTG GAGAGTTTTT GCAAGAGGGC CTGTAGTACC AGGGGGATTT 960
CTAACAGCTG CTGTCTGGAT GAGTTTTGCC CAAGCAGGAA TCAATCCTTT TGTCTGCATT1020
10 TTCTCAAACA GGGAGCTGAG GCGCTGTTC AGCACAACCC TTCTTTACTG CAGAAAATCC1080
AGGTTACCAA GGGAACCTTA CTGTGTTATA TGA

- (27) INFORMATION FOR SEQ ID NO:26:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 370 amino acids
- (B) TYPE: amino acid
  - (C) STRANDEDNESS:
  - (D) TOPOLOGY: not relevant
  - (ii) MOLECULE TYPE: protein
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:26:
- 20 Met Ala Asn Tyr Ser His Ala Ala Asp Asn Ile Leu Gln Asn Leu Ser 1 5 10 15
  - Pro Leu Thr Ala Phe Leu Lys Leu Thr Ser Leu Gly Phe Ile Ile Gly
    20 25 30
- Val Ser Val Val Gly Asn Leu Leu Ile Ser Ile Leu Leu Val Lys Asp

  35 40 45
  - Lys Thr Leu His Arg Ala Pro Tyr Tyr Phe Leu Leu Asp Leu Cys Cys 50 55 60
  - Ser Asp Ile Leu Arg Ser Ala Ile Cys Phe Pro Phe Val Phe Asn Ser 65 70 75 80
- Val Lys Asn Gly Ser Thr Trp Thr Tyr Gly Thr Leu Thr Cys Lys Val 85
  - Ile Ala Phe Leu Gly Val Leu Ser Cys Phe His Thr Ala Phe Met Leu

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		•		100			-);-	-++ · .	105				<i>:</i>	110		
-	Phe	: Cys	Ile 115	Ser	Val	Thr	Arg	Tyr 120	Leu	Ala	Ile	Ala	His		Arg	Phe
. 5	Tyr	Thr 130	Lys	Arg	Leu	Thr	Phe	Trp	Thr	Cys	Leu		Val	Ile	Cys	Met
	 Val		• •	T.e.u	Ser		715	 Mot	23.0	Db.		140		- ·		· .
	145			204		150	Ala	Met	AId	Pne	155	PIO	vaı	Leu	Asp	Val 160
.*	Gly	Thr	Tyr	Ser	Phe 165	Ile	Arg	Glu	Glu	Asp 170	Gln	Cys	Thr	Phe	Gln 175	His
10	Arg	Ser	Phe	Arg 180	Ala	Asn	Asp	Ser	Leu 185	Gly	Phe	Met	Leu	Leu 190	Leu	Ala
· · · · · · · · · · · · · · · · · · ·	Leu	Ile	Leu 195	Leu	Ala	Thr	Gln	Leu 200	Val	Tyr	Leu	Lys	Leu 205		Phe	Phe
15	Val	His 210	Asp	Arg	Arg	Lys	Met 215	Lys	Pro	Val	Gln	Phe 220	Val	Ala	Ala	Val
	Ser 225	Gln	Asn	Trp	Thr	Phe 230	His	Gly	Pro	Gly	Ala 235	Ser	Gly	Gln	Ala	Ala 240
	Ala	Asn	Trp	Leu	Ala 245	Gly	Phe	Gly	Arg	Gly 250	Pro	Thr	Pro	Pro	Thr 255	Leu
20	Leu	Gly	Ile	Arg 260	Gln	Asn	Ala	Asn	Thr 265	Thr	Gly	Arg	Arg	Arg 270	Leu	Leu
*, %	Val	Leu	Asp 275	Glu	Phe	Lys	Met	Glu 280	Lys	Arg	Ile	Ser	Arg 285	Met	Phe	Tyr
25	Ile	Met 290	Thr	Phe	Leu	Phe	Leu 295	Thr	Leu	Trp	Gly	Pro 300	Tyr	Leu	Val	Ala
	Cys 305	Tyr	Trp	Arg	Val	Phe 310	Ala	Arg	Gly	Pro	Val 315	Val	Pro	Gly	Gly	Phe 320
	Leu	Thr	Ala	Ala	Val 325	Trp	Met	Ser	Phe	Ala 330	Gln	Ala	Gly	Ile	Asn 335	Pro
30	Phe	Val	Cys	Ile 340	Phe	Ser	Asn	Arg	Glu 345	Leu	Arg	Arg	Cys	Phe 350	Ser	Thr
	Thr	Leu	Leu 355	Tyr	Cys	Arg	Lys	Ser 360	Arg	Leu	Pro	Arg	Glu 365		Tyr	Cys
35	Val		<del>-</del>					-				٠	303			
,,,		370	, ,			. •						- 131				

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- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1080 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:27:

ATGCAGGTCC CGAACAGCAC CGGCCCGGAC AACGCGACGC TGCAGATGCT GCGGAACCCG 60 GCGATCGCGG TGGCCCTGCC CGTGGTGTAC TCGCTGGTGG CGGCGGTCAG CATCCCGGGC 120 10 AACCTCTTCT CTCTGTGGGT GCTGTGCCGG CGCATGGGGC CCAGATCCCC GTCGGTCATC 180 TTCATGATCA ACCTGAGCGT CACGGACCTG ATGCTGGCCA GCGTGTTGCC TTTCCAAATC 240 TACTACCATT GCAACCGCCA CCACTGGGTA TTCGGGGTGC TGCTTTGCAA CGTGGTGACC 300 GTGGCCTTTT ACGCAAACAT GTATTCCAGC ATCCTCACCA TGACCTGTAT CAGCGTGGAG 360 CGCTTCCTGG GGGTCCTGTA CCCGCTCAGC TCCAAGCGCT GGCGCCGCCG TCGTTACGCG 420 15 GTGGCCGCGT GTGCAGGGAC CTGGCTGCTG CTCCTGACCG CCCTGTGCCC GCTGGCGCGC 480 ACCGATCTCA CCTACCCGGT GCACGCCCTG GGCATCATCA CCTGCTTCGA CGTCCTCAAG 540 TGGACGATGC TCCCCAGCGT GGCCATGTGG GCCGTGTTCC TCTTCACCAT CTTCATCCTG 600 CTGTTCCTCA TCCCGTTCGT GATCACCGTG GCTTGTTACA CGGCCACCAT CCTCAAGCTG 660 TTGCGCACGG AGGAGGCGCA CGGCCGGGAG CAGCGGAGGC GCGCGGTGGG CCTGGCCGCG 720 20 GTGGTCTTGC TGGCCTTTGT CACCTGCTTC GCCCCCAACA ACTTCGTGCT CCTGGCGCAC 780 ATCGTGAGCC GCCTGTTCTA CGGCAAGAGC TACTACCACG TGTACAAGCT CACGCTGTGT 840 CTCAGCTGCC TCAACAACTG TCTGGACCCG TTTGTTTATT ACTTTGCGTC CCGGGAATTC 900 CAGCTGCGCC TGCGGGAATA TTTGGGCTGC CGCCGGGTGC CCAGAGACAC CCTGGACACG 960 CGCCGCGAGA GCCTCTTCTC CGCCAGGACC ACGTCCGTGC GCTCCGAGGC CGGTGCGCAC1020 25 CCTGAAGGGA TGGAGGGAGC CACCAGGCCC GGCCTCCAGA GGCAGGAGAG TGTGTTCTGA1080

## (29) INFORMATION FOR SEQ ID NO:28:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 359 amino acids
  - (B) TYPE: amino acid
- (C) STRANDEDNESS:
  - (D) TOPOLOGY: not relevant

* Take		- 1		- 11	*			- ,55		0					7	
	(ii)	MO	LECU	LE T	YPE:	prot	tein				e. P.		• •	1. 1. v		12
				-					-	ý (			· · · ·			
	(xi)	SEC	QUENC	CE DI	ESCRI	PTIC	ON: S	SEQ 1	D NO	):28:	*					
: · · ·	Met	Gli	va]	l Pro	Asr	ı Ser	Thr	Glv	/ Pro	) Asc	Asn	Ala	ነ ጥኮን	i. Des	ı Glr	`. ∴Mat
*	1	· .			5				1	10		īš,		. He	15	Met
5	Leu	Arg	Asr	Pro	Ala	ile	Ala	Val	Ala	Leu	Pro	Val	. Val	Tvz	Ser	Leu
Y				20					25		• 1			30	4	
	Val	Ala	Ala	Val	Ser	Ile	Pro	Gly	Asn	Leu	Phe	Ser	Leu	Tre	Val	Leu
10			35	* **	1	1.0		40					45			
	Cys	Arg	Arg	Met	Gly	Pro	Arg	Ser	Pro	Ser	Val	Ile	Phe	Met	Ile	Asn
10		50				. :	55					60				
	Leu	Ser	Val	Thr	Asp	Leu	Met	Leu	Ala	Ser	Val	Leu	Pro	Phe	Gln	Ile
	65					70				. 111	75		¥	,		80
	Tyr	Tyr	His	Cys	Asn	Arg	His	His	Trp	Val	Phe	Gly	Val	Leu	Leu	Cvs
			e e	• 1,4	85		4			90		, O			95	-2-
15	Asn	Val	Val	Thr	Val	Ala	Phe	Tyr	Ala	Asn	Met	Tyr	Ser	Ser	Ile	Leu
				100					105	7	r ,	 		110		,,,,,,
	Thr	Met	Thr	Cys	Ile	Ser	Val	Glu	Arg	Phe	Leu	Gly	Val	Leu	Tyr	Pro
			115					120	-			•	125			
	Leu	Ser	Ser	Lys	Arg	Trp	Arg	Arg	Arg	Arg	Tyr	Ala	Val	Ala	Ala	Cvs
20		130			3.0		135					140				
	Ala	Gly	Thr	Trp	Leu	Leu	Leu	Leu	Thr	Ala	Leu	Cys	Pro	Leu	Ala	Arq
	145	*				150	(*)		7 - 2	s Light	155				. •	160
* * =	Thr	Asp	Leu	Thr	Tyr	Pro	Val	His	Ala	Leu	Gly	Île	Ile	Thr	Cys	Phe
					165	* .	1			170				1	175	× ·
25	Asp	Val	Leu	Lys	Trp	Thr	Met	Leu	Pro.	Ser	Val	Ala	Met	Tro	Ala	Val
				180					185					190		, , , ,
	Phe	Leu	Phe	Thr	Ile	Phe	Ile	Leu	Leu	Phe	Leu-	Ile	Pro	Phe	Val	Île
			195		<i>1</i> * * * * * * * * * * * * * * * * * * *		• . •	200					205			
	Thr	Val-	Ala	Cys	Tyr	Thr	Ala	Thr	Île	Leu	Lys	Leu	Leu	Arg	Thr	Glu
30		210	2		, :		215				*	220			-	
	Glu	Ala	His	Gly	Arg	Glu	Gln	Arg	Arg	Arg	Ala	Val	Gly	Leu	Ala	Ala
<b>(</b> )	225					230					235		×			240
	Val	Val	Leu	Leu	Ala	Phe	Val	Thr	Cys	Phe	Ala	Pro	Asn	Asn	Phe	Val
	- 1				245			• • • • •		250	Σ".		•		255	
35	Leu	Leu	Ala	His	Ile	Val	Ser	Arg	Leu	Phe	Tyr	Gly	Lys	Ser	Tyr	Tyr
w				260					265	· · · · · · · · · · · · · · · · · · ·		•	X.	270		-

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His Val Tyr Lys Leu Thr Leu Cys Leu Ser Cys Leu Asn Asn Cys Leu 275 280 285

- Asp Pro Phe Val Tyr Phe Ala Ser Arg Glu Phe Gln Leu Arg Leu 290 295 300
- Arg Glu Tyr Leu Gly Cys Arg Arg Val Pro Arg Asp Thr Leu Asp Thr 305
  - Arg Arg Glu Ser Leu Phe Ser Ala Arg Thr Thr Ser Val Arg Ser Glu 325 330 335
- Ala Gly Ala His Pro Glu Gly Met Glu Gly Ala Thr Arg Pro Gly Leu
  340 345 350

Gln Arg Gln Glu Ser Val Phe 355

#### (30) INFORMATION FOR SEQ ID NO:29:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1503 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- 20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:29:

ATGGAGCGTC CCTGGGAGGA CAGCCCAGGC CCGGAGGGGG CAGCTGAGGG CTCGCCTGTG 60

CCAGTCGCCG CCGGGGCGC CTCCGGTGCC GCGCCGAGTG GCACAGGCTG GCAGCCATGG 120

• GCTGAGTGCC CGGGACCCAA GGGGAGGGGG CAACTGCTGG CGACCGCCGG CCCTTTGCGT 180

CGCTGGCCCG CCCCCTCGCC TGCCAGCTCC AGCCCCGCCC CCGGAGCGGC GTCCGCTCAC 240

25 TCGGTTCAAG GCAGCGCGAC TGCGGGTGGC GCACCGACCAG GGCGCAGACC TTGGGGCGCG 300

CGGCCCATGG AGTCGGGGCT GCTGCGGCCG GCGCCGGTGA GCGAGGTCAT CGTCCTGCAT 360

TACAACTACA CCGGCAAGCT CCGCGGTGCG AGCTACCAGC CGGGTGCCGG CCTGCGCGCC 420

GACGCCGTGG TGTGCCTGGC GGTGTGCGC TTCATCGTGC TAGAGAAATCT AGCCGTGTTG 480

TTGGTGCTCG GACGCCACCC GCGCTTCCAC GCTCCCATGT TCCTGCTCCT GGGCAGCCTC 540

30 ACGTTGTCGG ATCTGCTGC AGGCGCCGCC TACGCCGCCA ACATCCTACT GTCGGGGCCG 600

CTCACGCTGA AACTGTCCCC CGCGCTCTGG TTCGCACGGG AGGGAGGCGT CTTCGTGGCA 660

CTCACTGCGT CCGTGCTGAG CCTCCTGGCC ATCGCGCTGG AGCGCAGCCT CACCATGGCG 720

TGGGGGGGGC CCGCGCCCGT CTCCAGTCGG GGGCGCACGC TGGCGATGGC AGCCGCGGCC 780

TGGGGCGTGT CGCTGCTCCT CGGGCTCCTG CCAGCGCTGG GCTGGAATTG CCTGGGTCGC 840

CTGGACGCTT GCTCCACTGT CTTGCCGCTC TACGCCAAGG CCTACGTGCT CTTCTGCGTG 900

CTCGCCTTCG TGGGCATCCT GGCCGCGATC TGTGCACTCT ACGCGCGCAT CTACTGCCAG 960

5 GTACGCGCCA ACGCGCGCG CCTGCCGGCA CGGCCCGGGA CTGCGGGGAC CACCTCGACC1020

CGGGCGCGTC GCAAGCCGCG CTCTCTGGCC TTGCTGCGCA CGCTCAGCGT GGTGCTCCTG1080

GCCTTTGTGG CATGTTGGGG CCCCCTCTTC CTGCTGCTGT TGCTCGACGT GGCGTGCCCG1140

GCGCGCACCT GTCCTGTACT CCTGCAGGCC GATCCCTTCC TGGGACTGC CATGGCCAAC1200

TCACTTCTGA ACCCCATCAT CTACACGCTC ACCAACCGCG ACCTGCGCCA CGCGCTCCTG1260

10 CGCCTGGTCT GCTGCGGACG CCACTCCTGC GGCAGAGACC CGAGTGGCTC CCAGCAGTCG1320

GCGAGCGCGG CTGAGGCTTC CGGGGGCCTG CGCCGCTGCC TGCCCCCGGG CCTTGATGGG1380

AGCTTCAGCG GCTCGGAGCG CTCATCGCCC CAGCGCGACG GGCTGGACAC CAGCGGCTCC1440

ACAGGCAGCC CCGGTGCACC CACAGCCGCC CGGACTCTGG TATCAGAACC GGCTGCAGAC1500

#### 15 (31) INFORMATION FOR SEQ ID NO:30:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 500 amino acids
    - (B) TYPE: amino acid
    - (C) STRANDEDNESS:
- 20 (D) TOPOLOGY: not relevant
  - (ii) MOLECULE TYPE: protein
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:30:

Met Glu Arg Pro Trp Glu Asp Ser Pro Gly Pro Glu Gly Ala Ala Glu

10
15

Gly Ser Pro Val Pro Val Ala Ala Gly Ala Arg Ser Gly Ala Ala Ala 20 25 30

Ser Gly Thr Gly Trp Gln Pro Trp Ala Glu Cys Pro Gly Pro Lys Gly
35 40 45

Arg Gly Gln Leu Leu Ala Thr Ala Gly Pro Leu Arg Arg Trp Pro Ala
50 55 60

Pro Ser Pro Ala Ser Ser Ser Pro Ala Pro Gly Ala Ala Ser Ala His
65 70 75 80

1.74	Sei	· Val	l Gln	Gly	Ser 85	Ala	Thr	Ala	Gly	Gly 90	Ala	Arg	Pro	Gly	Arg 95	Arg
	Pro	Trp	Gly	Ala 100		Pro	Met	Glu	Ser 105		Leu	Leu	Arg	Pro	Ala	Pro
5	Val	. Ser	Glu 115	Val	Ile	Val	Leu	His 120		Asn	Tyr	Thr	Gly 125		Leu	Arg
	Gly	Ala	Ser	Tyr	Gln	Pro	Gly 135		Gly	Leu	Arg	Ala 140	Asp	Ala	Val	Val
.10	Cys	Leu	Ala	Val	Cys	Ala 150	Phe	Ile	Val	Leu	Glu 155	Asn	Leu	Ala	Val	Leu 160
	Leu	Val	Leu	Gly	Arg 165	His	Pro	Arg	Phe	His 170	Ala	Pro	Met	Phe	Leu 175	Leu
	Leu	Gly	Ser	Leu 180	Thr	Leu	Ser	Asp	Leu 185	Leu	Ala	Gly	Ala	Ala 190	Tyr	Ala
15	Ala	Asn	Ile 195	Leu	Leu	Ser	Gly	Pro 200	Leu	Thr	Leu	Lys	Leu 205	Ser	Pro	Ala
	Leu	Trp 210	Phe	Ala	Arg	Glu	Gly 215		Val	Phe	Val	Ala 220	Leu	Thr	Ala	Ser
20	Val 225	Leu	Ser	Leu	Leu	Ala 230	Ile	Ala	Leu	Glu	Arg 235	Ser	Leu	Thr	Met	Ala 240
	Arg	Arg	Gly	Pro	Ala 245	Pro	Val	Ser	Ser	Arg 250	Gly	Arg	Thr	Leu	Ala 255	Met
	Ala	Ala	Ala	Ala 260	Trp	Gly	Val	Ser	Leu 265	Leu	Leu	Gly	Leu	Leu 270	Pro	Ala
25	Leu	Gly	Trp 275	Asn	Cys	Leu	Gly	Arg 280	Leu	Asp	Ala	Cys	Ser 285		Val	Leu
	Pro	Leu 290	Tyr	Ala	Lys	Ala	Tyr 295	Val	Leu	Phe	Cys	Val 300	Leu	Ala	Phe	Val
30	305		0.			310	*				Ala 315	·				320
				. =	325	; ;				330	-				335	8
25	· · · · · · · · · · · · · · · · · · ·		-	340			. ** *		345	*	Arg		. (4	350		
35			355		- )-			360			Val		365			* -
•	Leu	Phe	Leu	Leu	Leu	Leu	Leu	Asp	Val	Ala	Cys	Pro	Ala	Arg	Thr	Cys

370 375 380

Pro Val Leu Leu Gln Ala Asp Pro Phe Leu Gly Leu Ala Met Ala Asn 385 390 395 400

Ser Leu Leu Asn Pro Ile Ile Tyr Thr Leu Thr Asn Arg Asp Leu Arg

His Ala Leu Leu Arg Leu Val Cys Cys Gly Arg His Ser Cys Gly Arg
420 425 430

Asp Pro Ser Gly Ser Gln Gln Ser Ala Ser Ala Ala Glu Ala Ser Gly
435
440
445

Gly Leu Arg Arg Cys Leu Pro Pro Gly Leu Asp Gly Ser Phe Ser Gly
450
450
460

Ser Glu Arg Ser Ser Pro Gln Arg Asp Gly Leu Asp Thr Ser Gly Ser 465 470 475 480

Thr Gly Ser Pro Gly Ala Pro Thr Ala Ala Arg Thr Leu Val Ser Glu
485
490
495

Pro Ala Ala Asp

#### (32) INFORMATION FOR SEQ ID NO:31:

(i) SEQUENCE CHARACTERISTICS:

20 (A) LENGTH: 1029 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

25 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:31:

ATGCAAGCCG TCGACAATCT CACCTCTGCG CCTGGGAACA CCAGTCTGTG CACCAGAGAC 60
TACAAAATCA CCCAGGTCCT CTTCCCACTG CTCTACACTG TCCTGTTTTT TGTTGGACTT 120
ATCACAAATG GCCTGGCGAT GAGGATTTTC TTTCAAATCC GGAGTAAATC AAACTTTATT 180
ATTTTTCTTA AGAACACAGT CATTTCTGAT CTTCTCATGA TTCTGACTTT TCCATTCAAA 240
30 ATTCTTAGTG ATGCCAAACT GGGAACAGGA CCACTGAGAA CTTTTGTGTG TCAAGTTACC 300
TCCGTCATAT TTTATTTCAC AATGTATATC AGTATTTCAT TCCTGGGACT GATAACTATC 360
GATCGCTACC AGAAGACCAC CAGGCCATTT AAAACATCCA ACCCCAAAAA TCTCTTGGGG 420
GCTAAGATTC TCTCTGTTGT CATCTGGGCA TTCATGTTCT TACTCTCTTT GCCTAACATG 480

ATTCTGACCA ACAGGCAGCC GAGAGACAAG AATGTGAAGA AATGCTCTTT CCTTAAATCA 540
GAGTTCGGTC TAGTCTGGCA TGAAATAGTA AATTACATCT GTCAAGTCAT TTTCTGGATT 600
AATTTCTTAA TTGTTATTGT ATGTTATACA CTCATTACAA AAGAACTGTA CCGGTCATAC 660
GTAAGAACGA GGGGTGTAGG TAAAGTCCCC AGGAAAAAGG TGAACGTCAA AGTTTTCATT 720
5 ATCATTGCTG TATTCTTTAT TTGTTTTGTT CCTTTCCATT TTGCCCGAAT TCCTTACACC 780
CTGAGCCAAA CCCGGGATGT CTTTGACTGC ACTGCTGAAA ATACTCTGTT CTATGTGAAA 840
GAGAGCACTC TGTGGTTAAC TTCCTTAAAT GCATGCCTGG ATCCGTTCAT CTATTTTTC 900
CTTTGCAAGT CCTTCAGAAA TTCCTTGATA AGTATGCTGA AGTGCCCCAA TTCTGCAACA 960
TCTCTGTCCC AGGACAATAG GAAAAAAGAA CAGGATGGTG GTGACCCAAA TGAAGAGACT1020
10 CCAATGTAA

## (33) INFORMATION FOR SEQ ID NO:32:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 342 amino acids
  - (B) TYPE: amino acid
- 15 (C) STRANDEDNESS:
  - (D) TOPOLOGY: not relevant
  - (ii) MOLECULE TYPE: protein
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:32:
- Met Gln Ala Val Asp Asn Leu Thr Ser Ala Pro Gly Asn Thr Ser Leu

  10
  15
  - Cys Thr Arg Asp Tyr Lys Ile Thr Gln Val Leu Phe Pro Leu Leu Tyr
    20 25 30
  - Thr Val Leu Phe Phe Val Gly Leu Ile Thr Asn Gly Leu Ala Met Arg
    35 40 45
- 25 Ile Phe Phe Gln Ile Arg Ser Lys Ser Asn Phe Ile Ile Phe Leu Lys
  50 55 60
  - Asn Thr Val Ile Ser Asp Leu Leu Met Ile Leu Thr Phe Pro Phe Lys
    70 75 80
- Ile Leu Ser Asp Ala Lys Leu Gly Thr Gly Pro Leu Arg Thr Phe Val 85 90 95
  - Cys Gln Val Thr Ser Val Ile Phe Tyr Phe Thr Met Tyr Ile Ser Ile
    100 105 110
  - -Ser Phe Leu Gly Leu Ile Thr Ile Asp Arg Tyr Gln Lys Thr Thr Arg

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	Ser	. Val	. Val	. Ile	Trp	Δĺa	Phe	Met	Dhe	. T.a.	LTON	Con	. T			•
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	Ile	Leu	Thr	Asn	Arg	Gln	Pro	Ara	Asp	Lvs	Asn	Val	Tare	i. Litare	Cira	
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1	Phe	Leu	Lys	Ser	Glu	Phe	Gly	Leu	Val	Tro	His	Glu	Tla	Val	λen	Ф.
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10	Ile	Cys	Gln	Val	Ile	Phe	Trp	Ile	Asn	Phe	Leu	Ile	Val	Ile	Val	Cvs
			195	:				200					205			
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	Tyr	Thr	Leu	Ile	Thr	Lys	Glu	Leu	Tyr	Arg	Ser	Tyr	Val	Arg	Thr	Aro
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16	Gly	Val	Gly	Lys	Val	Pro	Arg	Lys	Lys	Val	Asn	Val	Lys	Val	Phe	Ile
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* . *	TTE	TTE	Ala	Val	Phe	Phe	Ile	Cys	Phe	Val	Pro	Phe	His	Phe	Ala	Arg
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	116	PLO	Tyr	260	Leu	ser	GIn	Thr		Asp	Val	Phe	Asp	Cys	Thr	Ala
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20 (24)	-:					-		-								
30 (34)	INFO	RMAT	ION	FOR	SEQ	ID N	0:33	:								
		ana								• •					•	
* ***	(1)				RACT				•	* -		•				
		(A)	LEN	GTH:	107	7 ba	se p	airs						•		
		(B)	TYP	E: n	ucle	ic a	cid									

- 35
- (C) STRANDEDNESS: single (D) TOPOLOGY: linear
  - (ii) MOLECULE TYPE: DNA (genomic)

### (xi) SEQUENCE DESCRIPTION: SEQ ID NO:33:

ATGTCGGTCT GCTACCGTCC CCCAGGGAAC GAGACACTGC TGAGCTGGAA GACTTCGCGG 60 GCCACAGGCA CAGCCTTCCT GCTGCTGGCG GCGCTGCTGG GGCTGCCTGG CAACGGCTTC 120 GTGGTGTGGA GCTTGGCGGG CTGGCGGCCT GCACGGGGC GACCGCTGGC GGCCACGCTT 180 5 GTGCTGCACC TGGCGCTGGC CGACGGCGCG GTGCTGCTGC TCACGCCGCT CTTTGTGGCC 240 TTCCTGACCC GGCAGGCCTG GCCGCTGGGC CAGGCGGGCT GCAAGGCGGT GTACTACGTG 300 TGCGCGCTCA GCATGTACGC CAGCGTGCTG CTCACCGGCC TGCTCAGCCT GCAGCGCTGC 360 CTCGCAGTCA CCCGCCCCTT CCTGGCGCCT CGGCTGCGCA GCCCGGCCCT GGCCCGCCGC 420 CTGCTGCTGG CGGTCTGGCT GGCCGCCCTG TTGCTCGCCG TCCCGGCCGC CGTCTACCGC 480 10 CACCTGTGGA GGGACCGCGT ATGCCAGCTG TGCCACCCGT CGCCGGTCCA CGCCGCCGCC 540 CACCTGAGCC TGGAGACTCT GACCGCTTTC GTGCTTCCTT TCGGGCTGAT GCTCGGCTGC 600 TACAGCGTGA CGCTGGCACG GCTGCGGGGC GCCCGCTGGG GCTCCGGGCG GCACGGGGCG 660 CGGGTGGGCC GGCTGGTGAG CGCCATCGTG CTTGCCTTCG GCTTGCTCTG GGCCCCCTAC 720 CACGCAGTCA ACCTTCTGCA GGCGGTCGCA GCGCTGGCTC CACCGGAAGG GGCCTTGGCG 780 15 AAGCTGGGCG GAGCCGGCCA GGCGGCGCGA GCGGGAACTA CGGCCTTGGC CTTCTTCAGT 840 TCTAGCGTCA ACCCGGTGCT CTACGTCTTC ACCGCTGGAG ATCTGCTGCC CCGGGCAGGT 900 CCCCGTTTCC TCACGCGGCT CTTCGAAGGC TCTGGGGAGG CCCGAGGGGG CGGCCGCTCT 960 AGGGAAGGGA CCATGGAGCT CCGAACTACC CCTCAGCTGA AAGTGGTGGG GCAGGGCCGC1020 GGCAATGGAG ACCCGGGGGG TGGGATGGAG AAGGACGGTC CGGAATGGGA CCTTTGA 1077 20 (35) INFORMATION FOR SEQ ID NO:34:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 358 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS:
- (D) TOPOLOGY: not relevant
  - (ii) MOLECULE TYPE: protein
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:34:
  - Met Ser Val Cys Tyr Arg Pro Pro Gly Asn Glu Thr Leu Leu Ser Trp

    5 10 15
- 30 Lys Thr Ser Arg Ala Thr Gly Thr Ala Phe Leu Leu Leu Ala Ala Leu

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		Let	ı Gly	/ Leu 35	Pro	Gly	Asn	Gly	Phe 40	· Val	Val	Trp	Ser	Leu 45	ı Ala	Gly	Trj
5		Arg	Pro	) Ala	Arg	Gly	Arg	Pro	Leu	Ala	Ala	Thr	Leu 60	Val	Leu	His	Let
-4		Ala 65	Leu	ı Ala	Asp	Gly	Ala 70	Val	Leu	Leu	ı Leu	Thr	Pro	Leu	Phe	Val	Ala 80
•		Phe	Leu	Thr	Arg	Gln 85	Ala	Trp	Pro	Leu	Gly 90	Gln	Ala	Gly	Cys	Lys 95	
10		Val	Tyr	Tyr	Val	Cys	Ala	Leu	Ser		Tyr	Ala	Ser	Val		Leu	Thr
		Gly	Leu	Leu	Ser		Gln	Arg		105 Leu		Val	Thr	Arg	110 Pro	Phe	Leu
		Ala	Pro	115 Arg		Arg	Ser	Pro	120 Ala	Leu	Ala	Arg	Arg	125 Leu	Leu	Leu	Ala
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20		His	Ala	Ala	Ala 180	His	Leu	Ser	Leu	Glu 185	Thr	Leu	Thr	Ala	Phe 190	Val	Leu
• • • •		Pro	Phe	Gly 195	Leu	Met	Leu	Gly	Cys 200	Tyr	Ser	Val	Thr	Leu 205	Ala	Arg	Leu
25		Arg	Gly 210	Ala	Arg	Trp	Gly	Ser 215	Gly	Arg	His	Gly	Ala 220	Arg	Val	Gly	Arg
	'	Leu 225	Val	Ser	Ala	Ile	Val 230	Leu	Ala	Pḥe	Gly	Leu 235	Leu	Trp	Ala	Pro	Tyr 240
		His	Ala	Val	Asn	Leu 245	Leu	Gln	Ala	Val	Ala 250	Ala	Leu	Ala	Pro	Pro	Glu
30		Gly	Ala	Leu	Ala 260	Lys	Leu	Gly	Gly	Ala 265	Gly	Gln	Ala	Ala	Arg 270	Ala	Gly
		Thr	Thr	Ala 275	Leu	Ala	Phe	Phe	Ser 280	Ser	Ser	Val	Asn	Pro 285	Val	Leu	Tyr
35		Val	Phe 290	Thr	Ala	Gly	Asp	Leu 295	Leu	Pro	Arg	Ala	Gly 300		Arg	Phe .	Leu
		Thr 305	Arg	Leu	Phe	Glu	Gly 310	Ser	Gly	Glu	Ala		٠.	Gly	Gly	Arg	
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Arg Glu Gly Thr Met Glu Leu Arg Thr Thr Pro Gln Leu Lys Val Val

Gly Gln Gly Arg Gly Asn Gly Asp Pro Gly Gly Gly Met Glu Lys Asp
340
345

-5--Gly Pro Glu Trp Asp Leu 355

## (36) INFORMATION FOR SEQ ID NO:35:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1005 base pairs
- 10 (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
  - (ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:35:

15 ATGCTGGGGA TCATGGCATG GAATGCAACT TGCAAAAACT GGCTGGCAGC AGAGGCTGCC 60 CTGGAAAAGT ACTACCTTTC CATTTTTAT GGGATTGAGT TCGTTGTGGG AGTCCTTGGA 120 AATACCATTG TTGTTTACGG CTACATCTTC TCTCTGAAGA ACTGGAACAG CAGTAATATT 180 TATCTCTTTA ACCTCTCTGT CTCTGACTTA GCTTTTCTGT GCACCCTCCC CATGCTGATA 240 AGGAGTTATG CCAATGGAAA CTGGATATAT GGAGACGTGC TCTGCATAAG CAACCGATAT 300 20 GTGCTTCATG CCAACCTCTA TACCAGCATT CTCTTTCTCA CTTTTATCAG CATAGATCGA 360 TACTTGATAA TTAAGTATCC TTTCCGAGAA CACCTTCTGC AAAAGAAAGA GTTTGCTATT 420 TTAATCTCCT TGGCCATTTG GGTTTTAGTA ACCTTAGAGT TACTACCCAT ACTTCCCCTT 480 ATAAATCCTG TTATAACTGA CAATGGCACC ACCTGTAATG ATTTTGCAAG TTCTGGAGAC 540 CCCAACTACA ACCTCATTTA CAGCATGTGT CTAACACTGT TGGGGTTCCT TATTCCTCTT 600 25 TTTGTGATGT GTTTCTTTTA TTACAAGATT GCTCTCTCC TAAAGCAGAG GAATAGGCAG 660 GTTGCTACTG CTCTGCCCCT TGAAAAGCCT CTCAACTTGG TCATCATGGC AGTGGTAATC 720 TTCTCTGTGC TTTTTACACC CTATCACGTC ATGCGGAATG TGAGGATCGC TTCACGCCTG 780 GGGAGTTGGA AGCAGTATCA GTGCACTCAG GTCGTCATCA ACTCCTTTTA CATTGTGACA 840 CGGCCTTTGG CCTTTCTGAA CAGTGTCATC AACCCTGTCT TCTATTTTCT TTTGGGAGAT 900 30 CACTTCAGGG ACATGCTGAT GAATCAACTG AGACACAACT TCAAATCCCT TACATCCTTT 960 AGCAGATGGG CTCATGAACT CCTACTTTCA TTCAGAGAAA AGTGA 1005

(37)	INFORMATION	FOR	SEO TO	NO.36.
,	TITL OIGHT TON	LOK	SEC ID	MO:30:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 334 amino acids
  - (B) TYPE: amino acid
- (C) STRANDEDNESS:
  - (D) TOPOLOGY: not relevant
  - (ii) MOLECULE TYPE: protein
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:36:
- Met Leu Gly Ile Met Ala Trp Asn Ala Thr Cys Lys Asn Trp Leu Ala

  1 5 10 15
  - Ala Glu Ala Ala Leu Glu Lys Tyr Tyr Leu Ser Ile Phe Tyr Gly Ile
    20 25 30
  - Glu Phe Val Val Gly Val Leu Gly Asn Thr Ile Val Val Tyr Gly Tyr
    35 40 45
- 15 Ile Phe Ser Leu Lys Asn Trp Asn Ser Ser Asn Ile Tyr Leu Phe Asn
  50 55 60
  - Leu Ser Val Ser Asp Leu Ala Phe Leu Cys Thr Leu Pro Met Leu Ile
    65 70 75
- Arg Ser Tyr Ala Asn Gly Asn Trp Ile Tyr Gly Asp Val Leu Cys Ile
  85 90 95
  - Ser Asn Arg Tyr Val Leu His Ala Asn Leu Tyr Thr Ser Ile Leu Phe 100 105 110
  - Leu Thr Phe Ile Ser Ile Asp Arg Tyr Leu Ile Ile Lys Tyr Pro Phe
    115 120 125
- 25 Arg Glu His Leu Leu Gln Lys Lys Glu Phe Ala Ile Leu Ile Ser Leu 130 135 140
  - Ala Ile Trp Val Leu Val Thr Leu Glu Leu Leu Pro Ile Leu Pro Leu 145 150 155 160
- Ile Asn Pro Val Ile Thr Asp Asn Gly Thr Thr Cys Asn Asp Phe Ala 165 170 175
  - Ser Ser Gly Asp Pro Asn Tyr Asn Leu Ile Tyr Ser Met Cys Leu Thr 180 185 190
  - Leu Leu Gly Phe Leu Ile Pro Leu Phe Val Met Cys Phe Phe Tyr Tyr
    195 200 205
- Lys Ile Ala Leu Phe Leu Lys Gln Arg Asn Arg Gln Val Ala Thr Ala
  210 215 220

Leu Pro Leu Glu Lys Pro Leu Asn Leu Val Ile Met Ala Val Val Ile
235 230 235 240

Phe Ser Val Leu Phe Thr Pro Tyr His Val Met Arg Asn Val Arg Ile
245 250 255

5 Ala Ser Arg Leu Gly Ser Trp Lys Gln Tyr Gln Cys Thr Gln Val Val , 260 265 270

Ile Asn Ser Phe Tyr Ile Val Thr Arg Pro Leu Ala Phe Leu Asn Ser 275 280 285

Val Ile Asn Pro Val Phe Tyr Phe Leu Leu Gly Asp His Phe Arg Asp
290
295
300

Met Leu Met Asn Gln Leu Arg His Asn Phe Lys Ser Leu Thr Ser Phe 305 310 315 320

Ser Arg Trp Ala His Glu Leu Leu Leu Ser Phe Arg Glu Lys 325 330

## 15 (38) INFORMATION FOR SEQ ID NO:37:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1296 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
- 20 (D) TOPOLOGY: linear
  - (ii) MOLECULE TYPE: DNA (genomic)
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:37:

ATGCAGGCGC TTAACATTAC CCCGGAGCAG TTCTCTCGGC TGCTGCGGGA CCACAACCTG 60

ACGCGGGAGC AGTTCATCGC TCTGTACCGG CTGCGACCGC TCGTCTACAC CCCAGAGCTG 120

25 CCGGGACGCG CCAAGCTGGC CCTCGTGCTC ACCGGCGTGC TCATCTTCGC CCTGGCGCTC 180

TTTGGCAATG CTCTGGTGTT CTACGTGGTG ACCCGCAGCA AGGCCATGCG CACCGTCACC 240

AACATCTTTA TCTGCTCCTT GGCGCTCAGT GACCTGCTCA TCACCTTCTT CTGCATTCCC 300

GTCACCATGC TCCAGAACAT TTCCGACAAC TGGCTGGGGG GTGCTTTCAT TTGCAAGATG 360

GTGCCATTTG TCCAGTCTAC CGCTGTTGTG ACAGAAATGC TCACTATGAC CTGCATTGCT 420

30 GTGGAAAGGC ACCAGGGACT TGTGCATCCT TTTAAAATGA AGTGGCAATA CACCAACCGA 480

AGGGCTTTCA CAATGCTAGG TGTGGTCTGG CTGGTGGCAG TCATCGTAGG ATCACCCATG 540

TGGCACGTGC AACAACTTGA GATCAAATAT GACTTCCTAT ATGAAAAGGA ACACATCTGC 600

TGCTTAGAAG AGTGGACCAG CCCTGTGCAC CAGGAAGATCT ACACCACCTT CATCCTTGTC 660

ATCCTCTCC TCCTGCCTCT TATGGTGATG CTTATTCTGT ACAGTAAAAT TGGTTATGAA 720

CTTTGGATAA AGAAAAGAGT TGGGGATGGT TCAGTGCTTC GAACTATTCA TGGAAAAGAA 780

ATGTCCAAAA TAGCCAGGAA GAAGAAACGA GCTGTCATTA TGATGGTGAC AGTGGTGGCT 840

CTCTTTGCTG TGTGCTGGGC ACCATTCCAT GTTGTCCATA TGATGATTGA ATACAGTAAT 900

5 TTTGAAAAGG AATATGATGA TGTCACAATC AAGATGATTT TTGCTATCGT GCAAATTATT 960

GGATTTTCCA ACTCCATCTG TAATCCCATT GTCTATGCAT TTATGAATGA AAACCTTCAAA1020

AAAAAATGTTT TGTCTGCAGT TTGTTATTGC ATAGTAAATA AAACCTTCTC TCCAGCACAA1080

AGGCATGGAA ATTCAGGAAT TACAATGATG CGGAAGAAAG CAAAGTTTTC CCTCAGAGAG1140

AATCCAGTGG AGGAAACCAA AGGAGAAGCA TTCAGTGATG GCAACATTGA AGTCAAATTG1200

10 TGTGAACAGA CAGAGGAGAA GAAAAAGCTC AAACGACATC TTGCTCTCTT TAGGTCTGAA1260

CTGGCTGAGA ATTCTCCTTT AGACAGTGGG CATTAA 1296

#### (39) INFORMATION FOR SEQ ID NO:38:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 431 amino acids
- 15 (B) TYPE: amino acid
  - (C) STRANDEDNESS:
    - (D) TOPOLOGY: not relevant
  - (ii) MOLECULE TYPE: protein
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:38:
- Met Gln Ala Leu Asn Ile Thr Pro Glu Gln Phe Ser Arg Leu Leu Arg

  1 5 10 15
  - Asp His Asn Leu Thr Arg Glu Gln Phe Ile Ala Leu Tyr Arg Leu Arg 20 25 30
- Pro Leu Val Tyr Thr Pro Glu Leu Pro Gly Arg Ala Lys Leu Ala Leu

  25 40 45
  - Val Leu Thr Gly Val Leu Ile Phe Ala Leu Ala Leu Phe Gly Asn Ala 50 55 60
  - Leu Val Phe Tyr Val Val Thr Arg Ser Lys Ala Met Arg Thr Val Thr 65 70 75 80
- Asn Ile Phe Ile Cys Ser Leu Ala Leu Ser Asp Leu Leu Ile Thr Phe
  85 90 95
  - Phe Cys Ile Pro Val Thr Met Leu Gln Asn Ile Ser Asp Asn Trp Leu

- 48

	100	
	Gly Gly Ala Phe Ile Cys Lys Met Val Pro Phe Val Gln Ser Thr	Ala
5	Val Val Thr Glu Met Leu Thr Met Thr Cys Ile Ala Val Glu Arg	His
	Gln Gly Leu Val His Pro Phe Lys Met Lys Trp Gln Tyr Thr Asn 145 150 155	Arg 160
	Arg Ala Phe Thr Met Leu Gly Val Val Trp Leu Val Ala Val Ile 165 170 175	
10	Gly Ser Pro Met Trp His Val Gln Gln Leu Glu Ile Lys Tyr Asp 180 185 190	Phe
	Leu Tyr Glu Lys Glu His Ile Cys Cys Leu Glu Glu Trp Thr Ser 195 200 205	Pro
15	Val His Gln Lys Ile Tyr Thr Thr Phe Ile Leu Val Ile Leu Phe 210 215 220	Leu
	Leu Pro Leu Met Val Met Leu Ile Leu Tyr Ser Lys Ile Gly Tyr (225 230 235	Glu 240
	Leu Trp Ile Lys Lys Arg Val Gly Asp Gly Ser Val Leu Arg Thr 1 245 250 255	
20	His Gly Lys Glu Met Ser Lys Ile Ala Arg Lys Lys Lys Arg Ala V 260 265 270	7al
* 0	Ile Met Met Val Thr Val Val Ala Leu Phe Ala Val Cys Trp Ala P 275 280 285	ro'
25	Phe His Val Val His Met Met Ile Glu Tyr Ser Asn Phe Glu Lys G 290 295 300	lu
	Tyr Asp Asp Val Thr Ile Lys Met Ile Phe Ala Ile Val Gln Ile I 305 310 315 3	le 20
	Gly Phe Ser Asn Ser Ile Cys Asn Pro Ile Val Tyr Ala Phe Met A 325 330 335	
30	Glu Asn Phe Lys Lys Asn Val Leu Ser Ala Val Cys Tyr Cys Ile Va 340 345 350	
* * *	Asn Lys Thr Phe Ser Pro Ala Gln Arg His Gly Asn Ser Gly Ile Th 355 360 365	*
35	Met Met Arg Lys Lys Ala Lys Phe Ser Leu Arg Glu Asn Pro Val Gl 370 375 380	4
	Glu Thr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Le 385 390 395 40	

Cys Glu Gln Thr Glu Glu Lys Lys Lys Leu Lys Arg His Leu Ala Leu 405 410 415

Phe Arg Ser Glu Leu Ala Glu Asn Ser Pro Leu Asp Ser Gly His
420 425 430

- 5 (40) INFORMATION FOR SEQ ID NO:39:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 24 base pairs
    - (B) TYPE: nucleic acid
    - (C) STRANDEDNESS: single
- 10 (D) TOPOLOGY: linear
  - (ii) MOLECULE TYPE: DNA (genomic)
    - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:39:

CTGTGTACAG CAGTTCGCAG AGTG

24

- (41) INFORMATION FOR SEQ ID NO:40:
- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 24 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- 20 (ii) MOLECULE TYPE: DNA (genomic)
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:40:

GAGTGCCAGG CAGAGCAGGT AGAC

24

- Julie - **31** 

- (42) INFORMATION FOR SEQ ID NO:41:
  - (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 31 base pairs
    - (B) TYPE: nucleic acid
    - (C) STRANDEDNESS: single
    - (D) TOPOLOGY: linear
    - (ii) MOLECULE TYPE: DNA (genomic)
- 30 (iv) ANTI-SENSE: NO
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:41:

CCCGAATTCC TGCTTGCTCC CAGCTTGGCC C

(43) INFORMATION FOR SEQ ID NO:42:

(i) SEQUENCE CHARACTERISTICS:	
(A) LENGTH: 32 base pairs	
(B) TYPE: nucleic acid	
(C) STRANDEDNESS: single	ر و المراجعة المراجعة والمراجعة المراجعة ا
(D) TOPOLOGY: linear	
(ii) MOLECULE TYPE, DV2	
(ii) MOLECULE TYPE: DNA (genomic)	الكراه أأوجرانها أنباها ولافيا بالامر العاشرة والمعطورات
(iv) ANTI-SENSE: YES	
	공의 교회생활회 나는 것이 있는 것이 말했다.
(xi) SEQUENCE DESCRIPTION: SEQ ID	NO.42.
TGTGGATCCT GCTGTCAAAG GTCCCATTCC GG	
	가게 : [조리나) 11 [제 : 11 : 11 : 11 : 12 : 12 : 13 : 13 : 13
10 (44) INFORMATION FOR SEQ ID NO:43:	
(i) SEQUENCE CHARACTERISTICS:	
(A) LENGTH: 20 base pairs	
(B) TYPE: nucleic acid	
(C) STRANDEDNESS: single	
(D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: DNA (genomic)	
ITE: DNA (genomic)	
(iv) ANTI-SENSE: NO	
(xi) SEQUENCE DESCRIPTION: SEQ ID 1	VO:43:
TCACAATGCT AGGTGTGGTC	
20 (4E) TYPOPUS	20
20 (45) INFORMATION FOR SEQ ID NO:44:	
(i) CROUDINGS GUILD	
(i) SEQUENCE CHARACTERISTICS:	
(A) LENGTH: 22 base pairs (B) TYPE: nucleic acid	
(C) STRANDEDNESS: single	
25 (D) TOPOLOGY: linear	
inear	
(ii) MOLECULE TYPE: DNA (genomic)	
(iv) ANTI-SENSE: YES	
	ر الله الله الله الله الله الله الله الل
(xi) SEQUENCE DESCRIPTION: SEQ ID NO	D:44:
TGCATAGACA ATGGGATTAC AG	
0 (46) INFORMATION TO	<b>22</b>
0 (46) INFORMATION FOR SEQ ID NO:45:	
(i) SECTIONOR OWNER	
(i) SEQUENCE CHARACTERISTICS:	
<ul><li>(A) LENGTH: 511 base pairs</li><li>(B) TYPE: nucleic acid</li></ul>	
'~' rise: nucleic acid	

	(D)	TOPOLOGY:	linear
• '	·~ /	TOTOMOGI.	T T T T C G T

- (ii) MOLECULE TYPE: DNA (genomic)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:45:
- TCACAATGCT AGGTGTGGTC TGGCTGGTGG CAGTCATCGT AGGATCACCC ATGTGGCACG 60

  5 TGCAACAACT TGAGATCAAA TATGACTTCC TATATGAAAA GGAACACATC TGCTGCTTAG 120

  AAGAGTGGAC CAGCCCTGTG CACCAGAAGA TCTACACCAC CTTCATCCTT GTCATCCTCT 180

  TCCTCCTGCC TCTTATGGTG ATGCTTATTC TGTACGTAAA ATTGGTTATG AACTTTGGAT 240

  AAAGAAAAGA GTTGGGGATG GTTCAGTGCT TCGAACTATT CATGGAAAAG AAATGTCCAA 300

  AATAGCCAGG AAGAAGAAAC GAGCTGTCAT TATGATGGTG ACAGTGGTGG CTCTCTTTGC 360

  10 TGTGTGCTGG GCACCATTCC ATGTTGTCCA TATGATGATT GAATACAGTA ATTTTGAAAA 420

  GGAATATGAT GATGTCACAA TCAAGATGAT TTTTGCTATC GTGCAAATTA TTGGATTTTC 480

  CAACTCCATC TGTAATCCCA TTGTCTATGC A 511
  - (47) INFORMATION FOR SEQ ID NO:46:
    - (i) SEQUENCE CHARACTERISTICS:

15

25

- (A) LENGTH: 21 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- 20 (iv) ANTI-SENSE: NO
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:46:

#### CTGCTTAGAA GAGTGGACCA G

21

- (48) INFORMATION FOR SEQ ID NO:47:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 22 base pairs
      - (B) TYPE: nucleic acid
      - (C) STRANDEDNESS: single
      - (D) TOPOLOGY: linear
  - (ii) MOLECULE TYPE: DNA (genomic)
- 30 (iv) ANTI-SENSE: NO

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···	.,	1311	/	, ,,	• 7

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:47:	
CTGTGCACCA GAAGATCTAC AC	22
(49) INFORMATION FOR SEQ ID NO:48:	
(12) INTOKRATION FOR SEQ ID, NO:48:	
(i) SEQUENCE CHARACTERISTICS:	
(A) LENGTH: 21 base pairs	المساوحة والحج على مسافحة المحسومين في المداح الفح على والم المراجع المواجع المحاجج المحاج
(B) TYPE: nucleic acid	
(C) STRANDEDNESS: single	
(D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: DNA (genomic)	
10 (iv) ANTI-SENSE: YES	
ANII DENOE: IES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:48:	
CAAGGATGAA GGTGGTGTAG A	21
(50) INFORMATION FOR SEQ ID NO:49:	
(i) SECUENCE CHARACTER	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 23 base pairs	
(B) TYPE: nucleic acid	
(C) STRANDEDNESS: single	
(D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: DNA (genomic)	
20 (iv) ANTI-SPACE, VEC	
20 (iv) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:49:	
1201. 210 ID NO:49:	
GTGTAGATCT TCTGGTGCAC AGG	33
	23
(51) INFORMATION FOR SEQ ID NO:50:	
(i) SEQUENCE CHARACTERISTICS:	
25 (A) LENGTH: 21 base pairs	
(B) TYPE: nucleic acid	
(C) STRANDEDNESS: single	
(D) TOPOLOGY: linear	
	. *
(ii) MOLECULE TYPE: DNA (genomic)	
30 (xi) SECUENCE DESCRIPTION CRO TRANS	
30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:50:	
GCAATGCAGG TCATAGTGAG C	
TOTAL TOTAL TOTAL CONTROLLER	21
(52) INFORMATION FOR SEC ID NO. 51	

(i) SEQUENCE CHARACTERISTICS:	
(A) LENGTH: 27 base pairs	
(B) TYPE: nucleic acid	
(C) STRANDEDNESS: single	
5 (D) TOPOLOGY: linear	
(b) TOPOLOGI: Timear	ter in the second secon
(ii) MOI POUT D MIND DATE (	
(ii) MOLECULE TYPE: DNA (genomic)	
(iii) inpominator	
(iii) HYPOTHETICAL: YES	
(iv) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID	NO:51:
10 TGGAGCATGG TGACGGGAAT GCAGAAG	
(53) INFORMATION FOR SEQ ID NO:52:	
(i) SEQUENCE CHARACTERISTICS:	
(A) LENGTH: 27 base pairs	
(B) TYPE: nucleic acid	
(C) STRANDEDNESS: single	
(D) TOPOLOGY: linear	
(b) TopoLogi: Timear	
(iii) NOT BOTT D. MIDD	
(ii) MOLECULE TYPE: DNA (genomic)	
(far) ham omen are	
(iv) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID	NO:52:
0	To
0 GTGATGAGCA GGTCACTGAG CGCCAAG	2
(54) INFORMATION FOR SEQ ID NO:53:	
*	
(i) SEQUENCE CHARACTERISTICS:	
(A) LENGTH: 23 base pairs	
(B) TYPE: nucleic acid	
5 (C) STRANDEDNESS: single	
(D) TOPOLOGY: linear	
1-,	8
(ii) MOLECULE TYPE: DNA (genomic)	
(11) FOLLECODE TIPE: DNA (genomic)	
(in) NYMT CONTO	
(iv) ANTI-SENSE: NO	
(xi) SEQUENCE DESCRIPTION: SEQ ID	NO:53:
	$Y_0$
OGCAATGCAGG CGCTTAACAT TAC	3

(55) INFORMATION FOR SEQ ID NO:54:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 22 base pairs

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- 54 -

(B) TYPE: nucleic acid	
(C) STRANDEDNESS: single	
(D) TOPOLOGY: linear	
10102001. IIIlear	
(ii) MOLECULE TYPE: DNA (genom	
, ITE: DNA (genom:	<b>LC)</b>
5 (iv) ANTI-SENSE: YES	ويوفأ بمدر تدليك والصاعد فأمدتها بسراء بأبياك الكارب وماريها
Table 188	
(xi) SEQUENCE DESCRIPTION: SEQ	المراجون والهيد المانك ومواريون الإسابي بوالي والموارك ويوارك
DESCRIPTION: SEQ	1D NO:54:
TTGGGTTACA ATCTGAAGGG CA	
CIGARGO CA	
(56) INFORMATION FOR SEQ ID NO:55:	
YOU THE STATE OF TOK BEQ. ID NO:55:	
(i) SEQUENCE CHARACTERISTICS:	
(A) LENGTH: 23 base pairs	
(B) TYPE: nucleic acid	
(C) STRANDEDNESS: single	
(D) TOPOLOGY: linear	
(37 TOPOLOGI: Tinear	
(ii) MOLECULE TYPE: DNA (genomi	ည့် ခေါင်းသည် ရှိရည်။ သမီးကျွေးကျောင်းရှိ
THE DIA (GENOMI	C)
15 (iv) ANTI-SENSE: NO	
January No.	
(xi) SEQUENCE DESCRIPTION: SEQ	
SEQ.	LD NO:55:
ACTCCGTGTC CAGCAGGACT CTG	
	23
(57) INFORMATION FOR SEQ ID NO:56:	
(i) SEQUENCE CHARACTERISTICS:	
20 (A) LENGTH: 24 base pairs	
(B) TYPE: nucleic acid	
(C) STRANDEDNESS: single	
(D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: DNA (genomic	
,	
25 (iv) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ I	D NO.56.
TGCGTGTTCC TGGACCCTCA CGTG	
	24
(58) INFORMATION FOR SEQ ID NO:57:	
224 10 10:37:	*

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 29 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

,		- 55 -				
(ii	) MOLECULE TYPE: D	NA (genomic)		0	* *	
(iv	) ANTI-SENSE: NO				* * * * * * * * * * * * * * * * * * *	o 1
•	, ruit pense. No	*	The second of th			٠.
	· · · · · · · · · · · · · · · · · · ·			(1)		٠.
(xi	) SEQUENCE DESCRIP	TION: SEQ ID	NO:57:			
CAGGCCT	IGG ATTTTAATGT CAG	GGA <b>TGG</b>				2
5 (59) IN	FORMATION FOR SEQ	ID NO:58:		•		
	*				- ()	
(1)	SEQUENCE CHARACTI	ERISTICS:				
*	(A) LENGTH: 27 ]	base pairs	*		<u></u>	
	(B) TYPE: nucle:	ic acid	•			
10	(C) STRANDEDNESS (D) TOPOLOGY: 1:	s: single	- ()		-1-	*
5-2	(D) TOPOLOGI: I	inear		* * .		٠.
(ii)	MOLECULE TYPE: DI	VA (cenomic)		**	* *	
		(g. nomic)			, in the second	
. (iv)	ANTI-SENSE: YES			• •		
•						
•						
(xi)	SEQUENCE DESCRIPT	TION: SEQ ID	NO:58:			
GGAGAGTC	AG CTCTGAAAGA ATTC	CAGG			0	2
15 (22)			*	* **	=	
15 (60) INF	ORMATION FOR SEQ I	D NO:59:		- "		
**************************************	SEQUENCE CHARACTE	D.T.O.T.C.		0.1		
(1)	(A) LENGTH: 27 b					
	(B) TYPE: nuclei	ase pairs	1			
	(C) STRANDEDNESS					
20	(D) TOPOLOGY: li	singre		30		
	(= ) 10102001. 11	iicar			*	
(ii)	MOLECULE TYPE: DN	A (genomic)		* s		
(iv)	ANTI-SENSE: NO	*	. •		• .	
	0			.,	4.	
		•	* * * * * * * * * * * * * * * * * * * *			•
(xi)	SEQUENCE DESCRIPT:	ION: SEQ ID I	NO:59:	. X		٠.
TGATGTGAT	CG CCAGATACTA ATAG	CAC	*	r 👉 " į į		27
25 (61) INFO	RMATION FOR SEQ II	D NO:60:	***	V****		
/23	OPONE		*,		** * * * * * * * * * * * * * * * * * *	
(1)	SEQUENCE CHARACTER		(*)			
	(A) LENGTH: 27 ba	ase pairs		*		
	(B) TYPE: nucleio	c acid				

(ii) MOLECULE TYPE: DNA (genomic)

(D) TOPOLOGY: linear

(C) STRANDEDNESS: single

(iv) ANTI-SENSE: YES

30

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:60:	-
CCTGATTCAT TTAGGTGAGA TTGAGAC	
	27
(62) INFORMATION FOR SEQ ID NO:61:	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 22 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: single  (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: DNA (genomic)	
10 (iv) ANTI-SENSE: NO	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:61:	
GACAGGTACC TTGCCATCAA G	1
(63) INFORMATION FOR SEQ ID NO:62:	
(i) SEQUENCE CHARACTERISTICS:  15 (A) LENGTH: 22 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single	
(D) TOPOLOGY: linear	
(D) TOPOLOGY: linear  (ii) MOLECULE TYPE: DNA (genomic)	
(D) TOPOLOGY: linear	
(D) TOPOLOGY: linear  (ii) MOLECULE TYPE: DNA (genomic)	
(D) TOPOLOGY: linear  (ii) MOLECULE TYPE: DNA (genomic)  20 (iv) ANTI-SENSE: YES  (xi) SEQUENCE DESCRIPTION: SEQ ID NO:62:  CTGCACAATG CCAGTGATAA GG	`````````````````````````````````````
(D) TOPOLOGY: linear  (ii) MOLECULE TYPE: DNA (genomic)  20 (iv) ANTI-SENSE: YES  (xi) SEQUENCE DESCRIPTION: SEQ ID NO:62:  CTGCACAATG CCAGTGATAA GG	
(D) TOPOLOGY: linear  (ii) MOLECULE TYPE: DNA (genomic)  20 (iv) ANTI-SENSE: YES  (xi) SEQUENCE DESCRIPTION: SEQ ID NO:62:  CTGCACAATG CCAGTGATAA GG	2
(D) TOPOLOGY: linear  (ii) MOLECULE TYPE: DNA (genomic)  20 (iv) ANTI-SENSE: YES  (xi) SEQUENCE DESCRIPTION: SEQ ID NO:62:  CTGCACAATG CCAGTGATAA GG  (64) INFORMATION FOR SEQ ID NO:63:  (i) SEQUENCE CHARACTERISTICS:  (a) LENGTH: 27 base pairs  (b) TYPE: nucleic acid  (c) STRANDEDNESS: single	2
(D) TOPOLOGY: linear  (ii) MOLECULE TYPE: DNA (genomic)  20 (iv) ANTI-SENSE: YES  (xi) SEQUENCE DESCRIPTION: SEQ ID NO:62:  CTGCACAATG CCAGTGATAA GG  (64) INFORMATION FOR SEQ ID NO:63:  (i) SEQUENCE CHARACTERISTICS:  (i) SEQUENCE CHARACTERISTICS:  (B) TYPE: nucleic acid  (C) STRANDEDNESS: single  (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: DNA (genomic)  20 (iv) ANTI-SENSE: YES  (xi) SEQUENCE DESCRIPTION: SEQ ID NO:62:  CTGCACAATG CCAGTGATAA GG  (64) INFORMATION FOR SEQ ID NO:63:  (i) SEQUENCE CHARACTERISTICS:  (i) SEQUENCE CHARACTERISTICS:  (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear  (ii) MOLECULE TYPE: DNA (genomic)	

(65)	INFORMATION	FOR	SEO: TD	NO . 64 .

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 27 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (iv) ANTI-SENSE: YES
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:64:

#### 10 AGACCAGCCA GGGCACGCTG AAGAGTG

27

- (66) INFORMATION FOR SEQ ID NO:65:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 32 base pairs
    - (B) TYPE: nucleic acid
    - (C) STRANDEDNESS: single
      - (D) TOPOLOGY: linear
  - (ii) MOLECULE TYPE: DNA (genomic)
  - (iv) ANTI-SENSE: NO

15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:65:

#### 20 GATCAAGCTT CCATCCTACT GAAACCATGG TC

. 32

- (67) INFORMATION FOR SEQ ID NO:66:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 35 base pairs
    - (B) TYPE: nucleic acid
- 25 (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
  - (ii) MOLECULE TYPE: DNA (genomic)
  - (iv) ANTI-SENSE: YES
    - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:66:

### 30 GATCAGATCT CAGTTCCAAT ATTCACACCA CCGTC

35

- (68) INFORMATION FOR SEQ ID NO:67:
  - (i) SEQUENCE CHARACTERISTICS:

- 58 -

(A) LENGTH: 22 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear
5 (ii) MOLECULE TYPE: DNA (genomic)
(iv) ANTI-SENSE: NO
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:67:
CTGGTGTGCT CCATGGCATC CC
(69) INFORMATION FOR SEQ ID NO:68:
10 (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 22 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: single  (D) TOPOLOGY: linear
15 (ii) MOLECULE TYPE: DNA (genomic)
(iv) ANTI-SENSE: YES
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:68:
GTAAGCCTCC CAGAACGAGA GG
(70) INFORMATION FOR SEQ ID NO:69:
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 24 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: single  (D) TOPOLOGY: linear
25 (ii) MOLECULE TYPE: DNA (genomic)
(iv) ANTI-SENSE: NO  (xi) SEQUENCE DESCRIPTION: SEQ ID NO:69:
CAGCGCAGGG TGAAGCCTGA GAGC
(71) INFORMATION FOR SEQ ID NO:70:
30 (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 24 base pairs  (B) TYPE: nucleic acid

(C) STRANDEDNESS: single (D) TOPOLOGY: linear

(iv) ANTI-SENSE: NO

	1333	NOT BOTH B			· · · · ·	•	,	
	(11)	MOLECULE	TYPE: DN	A (genomi	c)	,		
	(iv)	ANTI-SEN	SE: YES		*		•	
	(xi)	SEQUENCE	DESCRIPT	ION: SEQ	ID NO:70:	-1		
GGC	ACCTGC	T GTGACC	IGTG CAGG					2
5 (72	) INFO	RMATION	FOR SEQ I	D NO:71:			-	
10	(i)	(A) LENG (B) TYPI (C) STRI	CHARACTE GTH: 22 b E: nuclei ANDEDNESS DLOGY: li	ase pairs c acid : single				
	(ii)	MOLECULE	TYPE: DN	A (genomic	:)			* *
. 2	(iv) :	ANTI-SENS	E: NO			* , * , * , * ;		
•	(xi)	SEQUENCE	DESCRIPTI	ON: SEQ I	D NO:71:			
GTC	CTGCCA	CTTCGAGA	CAT GG			*		22
15 (73)	INFO	RMATION F	OR SEQ ID	NO:72:		*	*	*
20	(i) s	(A) LENG (B) TYPE (C) STRA	CHARACTER TH: 23 ba : nucleic NDEDNESS: LOGY: lin	se pairs acid single				*
	(ii) <sub>,</sub> M	OLECULE	TYPE: DNA	(genomic	)			
		NTI-SENS		ON: SEQ II	D NO.72.			
GAAA		TGCCCTT			_ 1.0.,2.	*	<b>-</b> ,	23
25 (74)	INFOR	MATION FO	OR SEQ ID	NO:73:	a.			
30		(A) LENGT (B) TYPE: (C) STRAM	CHARACTER CH: 26 bas nucleic IDEDNESS:	se pairs acid single				*
	(ii) M	OLECULE 1	YPE: DNA	(genomic)				·

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:73:

CCAACACCAG CATCCATGGC ATCAAG

26

(75) INFORMATION FOR SEQ ID NO:74:

(i) SEQUENCE CHARACTERISTICS:

5

- (A) LENGTH: 27 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

10 (iv) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:74:

GGAGAGTCAG CTCTGAAAGA ATTCAGG

27

#### WORLD INTELLECTUAL PROPERTY ORGANIZATION International Bureau



#### INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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C12N 15/12, C07K 14/72			

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60/156,634

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09/417,044

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1 October 1999 (01.10.99)

12 October 1999 (12.10.99)

12 October 1999 (12.10.99)

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With international search report.

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(54) Title: HUMAN ORPHAN G PROTEIN-COUPLED RECEPTORS

#### (57) Abstract

The invention disclosed in this patent document relates to transmembrane receptors, more particularly to endogenous, human orphan G protein-coupled receptors.

## FOR THE PURPOSES OF INFORMATION ONLY

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	Cameroon		KP ,	Democratic People's		Z	New Zealand		٠.	
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CZ			KZ	Kazakstan		0	Romania			
DE	Czech Republic	0	LC	Saint Lucia	R	U ·	Russian Federation	15	Ä . • .	* *
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EE	Estonia		LR ·	Liberia	S	G	Singapore			

onal Application No PCT/US 99/23687

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IPC 7 C12N15/12 C07K14/72

According to International Patent Classification (IPC) or to both national classification and IPC

#### B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  $IPC \ 7 \ C12N \ C07K$ 

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT						
Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.				
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Further documents are listed in the continuation of box C.	Patent family members are listed in annex.
*A" document defining the general state of the art which is not considered to be of particular relevance  "E" earlier document but published on or after the international filing date  "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)  "O" document referring to an oral disclosure, use, exhibition or other means  "P" document published prior to the International filing date but later than the priority date claimed	<ul> <li>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</li> <li>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</li> <li>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</li> <li>"&amp;" document member of the same patent family</li> </ul>
Date of the actual completion of the international search  14 July 2000	Date of mailing of the international search report  0 2. 08. 00
Name and mailing address of the ISA  European Patent Office, P.B. 5818 Patentlaan 2  NL – 2280 HV Rijswijk  Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  Fax: (+31-70) 340-3016	Authorized officer  Mandl, B

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X	WO 98 50549 A (HUMAN GENOME SCIENCES INC; LI YI (US); RUBEN STEVEN M (US)) 12 November 1998 (1998-11-12) SEQ.IDs. 1 and 2	61-64
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BXI	Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)
÷.,.	
i nis int	emational Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:
1	Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
يا ۾ بنائ	a particularly and particular section of the sectio
10-1	이 이 방으로 되는 사이 경기를 들어 보니는 사람들은 회사를 하다고 있는 것을 하는 것이 살아 있다.
	하다 내 내 내 내 내 내 내 내 내 내 내 내 내 내 내 내 내 내 내
	[12] [4] 사고 다른 이토리는 이름이 가고 말았다. 주민이 모양하는 생각이 다른 다른
2.	Claims Nos.:
	because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
	지근 소장 교회의 회교 하는 일을 모으고 전혀 하셨다면 그리고 뭐 없지지 않아 마다라고
	그렇게 있었다. 그리면 하는데 그리 가장하다고 중 한다리 만든 강하나, 하나, 것이 나를 다 하다니
	영어는 이 경영 등은 독일을 때 그 일 속인하다. 그리고 경영한 역원으로 한 보다면서 전하다면 살았습니.
	설명이는 이 경험을 제되고 되는 하다. 이번 점에 들어 되어 하지 않아 있습니다면 하는 수 있습니다.
3	Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
	The state of the s
Box II	Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)
	Observations where unity of invention is facking (Continuation of Item 2 of first sheet)
This Inte	ernational Searching Authority found multiple inventions in this international application, as follows:
	see additional sheet
	See additional Sheet
	합니다. 그는 그 아이들들은 아무리가 그리고 하는 이 하는 이 사고를 보고 있다. 그리고 있다.
	공기 의 기본 전 교육적인 이 그는 영화가를 살아 하셨습니다. 얼마야 되어 그리고 없는 없다.
<u></u> -	레이노(영리는 이 뭐들다.)하는 사람들에게 살아가면 이렇게 보다고 모든다.)
1. X	As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
	Social due Calific.
2	As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
	보이는 이미를 하고 말이 하다는 나와 아름답니다는 어린다리 하다를 걸으로 살랐다.
3.	As only some of the required additional search fees were timely paid by the applicant, this International Search Report
لاا	covers only those claims for which fees were paid, specifically claims Nos.:
. (* )	
- •	
4.	No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is
	restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
1 1	
Remark	on Protest The additional search fees were accompanied by the applicant's protest.
ــــــــــــــــــــــــــــــــــــــ	No protest accompanied the payment of additional search fees.

#### FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

#### 1. Claims: 1-4

Human G protein-coupled receptor as characterized by SEQ.ID.2, a cDNA encoding said receptor as characterized by SEQ.ID.1, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 2. Claims: 5-8

Human G protein-coupled receptor as characterized by SEQ.ID.4, a cDNA encoding said receptor as characterized by SEQ.ID.3, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 3. Claims: 9-12

Human G protein-coupled receptor as characterized by SEQ.ID.6, a cDNA encoding said receptor as characterized by SEQ.ID.5, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 4. Claims: 13-16

Human G protein-coupled receptor as characterized by SEQ.ID.8, a cDNA encoding said receptor as characterized by SEQ.ID.7, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 5. Claims: 17-20

Human G protein-coupled receptor as characterized by SEQ.ID.10, a cDNA encoding said receptor as characterized by SEQ.ID.9, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 6. Claims: 21-24

Human G protein-coupled receptor as characterized by SEQ.ID.12, a cDNA encoding said receptor as characterized by SEQ.ID.11, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 7. Claims: 25-28

Human G protein-coupled receptor as characterized by SEQ.ID.14, a cDNA encoding said receptor as characterized by

#### FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

SEQ.ID.13, a plasmid comprising said cDNA, and a host cell-comprising said plasmid.

#### 8. Claims: 29-32

Human G protein-coupled receptor as characterized by SEQ.ID.16, a cDNA encoding said receptor as characterized by SEQ.ID.15, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 9. Claims: 33-36

Human G protein-coupled receptor as characterized by SEQ.ID.18, a cDNA encoding said receptor as characterized by SEQ.ID.17, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 10. Claims: 37-40

Human G protein-coupled receptor as characterized by SEQ.ID.20, a cDNA encoding said receptor as characterized by SEQ.ID.19, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 11. Claims: 41-44

Human G protein-coupled receptor as characterized by SEQ.ID.22, a cDNA encoding said receptor as characterized by SEQ.ID.21, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 12. Claims: 45-48

Human G protein-coupled receptor as characterized by SEQ.ID.24, a cDNA encoding said receptor as characterized by SEQ.ID.23, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 13. Claims: 49-52

Human G protein-coupled receptor as characterized by SEQ.ID.26, a cDNA encoding said receptor as characterized by SEQ.ID.25, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 14. Claims: 53-56

#### FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

SEQ.ID.28, a cDNA encoding said receptor as characterized by SEQ.ID.27, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 15. Claims: 57-60

Human G protein-coupled receptor as characterized by SEQ.ID.30, a cDNA encoding said receptor as characterized by SEQ.ID.29, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 16. Claims: 61-64

Human G protein-coupled receptor as characterized by SEQ.ID.32, a cDNA encoding said receptor as characterized by SEQ.ID.31, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 17. Claims: 65-68

Human G protein-coupled receptor as characterized by SEQ.ID.34, a cDNA encoding said receptor as characterized by SEQ.ID.33, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 18. Claims: 69-72

Human G protein-coupled receptor as characterized by SEQ.ID.36, a cDNA encoding said receptor as characterized by SEQ.ID.35, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

#### 19. Claims: 73-76

Human G protein-coupled receptor as characterized by SEQ.ID.38, a cDNA encoding said receptor as characterized by SEQ.ID.37, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

information on patent family members

Inta onal Application No PCT/US 99/23687

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